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=> s (perfluoropolyether or polyether or polyoxyalkylene ether) and (gas? (2a) hydrogen or reduc? or hydrogenation) and (group viii or palladium or platinum or rhodium or ruthenium or pd or pt or rh or ru) and (support? (5a) fluoride)

9 FILES SEARCHED...

14 FILES SEARCHED...

22 FILES SEARCHED...

28 FILES SEARCHED...

36 FILES SEARCHED...

41 FILES SEARCHED...

49 FILES SEARCHED...

52 FILES SEARCHED...

58 FILES SEARCHED...

62 FILES SEARCHED...

64 FILES SEARCHED...

67 FILES SEARCHED...

75 FILES SEARCHED...

L1 87 (PERFLUOROPOLYETHER OR POLYETHER OR POLYOXYALKYLENE ETHER) AND  
(GAS? (2A) HYDROGEN OR REDUC? OR HYDROGENATION) AND (GROUP VIII  
OR PALLADIUM OR PLATINUM OR RHODIUM OR RUTHENIUM OR PD OR PT OR  
RH OR RU) AND (SUPPORT? (5A) FLUORIDE)

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L2 82 DUP REM L1 (5 DUPLICATES REMOVED)

=> d 12 1-82 ti

L2 ANSWER 1 OF 82 PROMT COPYRIGHT 2004 Gale Group on STN

TI Trade name directory.

L2 ANSWER 2 OF 82 CAPLUS COPYRIGHT 2004 ACS on STN

TI Process for the preparation of perfluoropolyethers having aldehyde,  
alcohol, and amine end groups by catalytic reduction

L2 ANSWER 3 OF 82 EUROPATFULL COPYRIGHT 2004 WILA on STN

TIEN Azeotropic compositions comprising 1,1,1,2,3,3-heptafluoropropane and  
processes using said compositions.

L2 ANSWER 4 OF 82 EUROPATFULL COPYRIGHT 2004 WILA on STN

TIEN PROTON-CONDUCTIVE POLYMER FILM AND PROCESS FOR PRODUCING THE SAME.

L2 ANSWER 5 OF 82 EUROPATFULL COPYRIGHT 2004 WILA on STN

TIEN PROCESSES FOR THE PRODUCTION OF HEXAFLUOROPROPENE AND OPTIONAL OTHER  
HALOGENATED HYDROCARBONS CONTAINING FLUORINE.

L2 ANSWER 6 OF 82 EUROPATFULL COPYRIGHT 2004 WILA on STN

TIEN PROCESSES FOR THE MANUFACTURE OF 1,1,1,3,3-PENTAFLUOROPROPENE,  
2-CHLORO-PENTAFLUOROPROPENE.

L2 ANSWER 7 OF 82 PCTFULL COPYRIGHT 2004 Univentio on STN

10/630,698

TIEN ELECTROSTATIC PROCESSING OF ELECTROCHEMICAL DEVICE COMPONENTS  
TIFR TRAITEMENT ELECTROSTATIQUE DE COMPOSANTS DE DISPOSITIFS ELECTROCHIMIQUES

L2 ANSWER 8 OF 82 USPATFULL on STN  
TI Fuel cell, fuel cell generator, and equipment using the same

L2 ANSWER 9 OF 82 USPATFULL on STN  
TI Proton-conductive polymer film and process for producing the same

L2 ANSWER 10 OF 82 USPATFULL on STN  
TI Reagents and methods for library synthesis and screening

L2 ANSWER 11 OF 82 PROMT COPYRIGHT 2004 Gale Group on STN

TI Trade name directory. (A-O).

L2 ANSWER 12 OF 82 EUROPATFULL COPYRIGHT 2004 WILA on STN  
TIEN Polymerization of cyclic ethers using heterogeneous catalysts.  
TIEN Polymerization of cyclic ethers using heterogeneous catalysts.

L2 ANSWER 13 OF 82 USPATFULL on STN DUPLICATE 1  
TI Production of 1,2-dihydro and 2,2-dihydro hexafluoropropanes and  
azeotropes thereof with HF

L2 ANSWER 14 OF 82 USPATFULL on STN DUPLICATE 2  
TI Process for the manufacture of 1,1,1,3,3-pentafluoropropene,  
2-chloro-pentafluoropropene and compositions comprising saturated  
derivatives

L2 ANSWER 15 OF 82 USPATFULL on STN DUPLICATE 3  
TI PROCESSES FOR THE MANUFACTURE OF 1,1,1,3,3- PENTAFLUOROPROPENE,  
2-CHLORO-PENTAFLUOROPROPENE AND COMPOSITIONS COMPRISING SATURATED  
DERIVATIVES THEREOF

L2 ANSWER 16 OF 82 EUROPATFULL COPYRIGHT 2004 WILA on STN  
TIEN Fuel cell, fuel cell generator, and equipment using the same.

L2 ANSWER 17 OF 82 USPATFULL on STN  
TI Fuel cell, fuel cell generator, and equipment using the same

L2 ANSWER 18 OF 82 USPATFULL on STN  
TI Interfacially polymerized, bipiperidine-polyamide membranes for reverse  
osmosis and/or nanofiltration and process for making the same

L2 ANSWER 19 OF 82 USPATFULL on STN  
TI Production of 1,2-dihydro and 2,2-dihydro hexafluoropropanes and  
azeotropes thereof with HF

L2 ANSWER 20 OF 82 USPATFULL on STN DUPLICATE 4  
TI Fuel cell with monolithic flow field-bipolar plate assembly and method  
for making and cooling a fuel cell stack

L2 ANSWER 21 OF 82 USPATFULL on STN DUPLICATE 5  
TI Gas diffusion electrode with nanosized pores and method for making same

L2 ANSWER 22 OF 82 PCTFULL COPYRIGHT 2004 Univentio on STN  
TIEN GAS DIFFUSION ELECTRODE WITH NANOSIZED PORES AND METHOD FOR MAKING SAME  
TIFR ELECTRODE DE DIFFUSION GAZEUSE A PORES DE TAILLE NANOMETRIQUE ET PROCEDE  
POUR LA FABRICATION D'UNE TELLE ELECTRODE

L2 ANSWER 23 OF 82 PCTFULL COPYRIGHT 2004 Univentio on STN  
TIEN FUEL CELL WITH MONOLITHIC FLOW FIELD-BIPOLAR PLATE ASSEMBLY AND METHOD

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- TIFR FOR MAKING AND COOLING A FUEL CELL STACK  
PILE A COMBUSTIBLE A ASSEMBLAGE DE PLAQUES A CHAMP BIPOLAIRE ET  
ECOULEMENT MONOLITHIQUE, ET PROCEDE DE FABRICATION ET DE REFROIDISSEMENT  
D'UN EMPILEMENT DE PILES A COMBUSTIBLE
- L2 ANSWER 24 OF 82 PCTFULL COPYRIGHT 2004 Univentio on STN  
TIEN ELECTRONICALLY CONDUCTING FUEL CELL COMPONENT WITH DIRECTLY BONDED  
LAYERS AND METHOD FOR MAKING SAME
- TIFR COMPOSANT DE PILE A COMBUSTIBLE CONDUCTEUR SUR LE PLAN ELECTRONIQUE DOTE  
DE COUCHES DIRECTEMENT LIEES ET PROCEDE DE FABRICATION CORRESPONDANT
- L2 ANSWER 25 OF 82 USPATFULL on STN  
TI Electronically conducting fuel cell component with directly bonded  
layers and method for making same
- L2 ANSWER 26 OF 82 USPATFULL on STN  
TI Processes for the production of hexafluoropropene and optionally other  
halogenated hydrocarbons containing fluorine
- L2 ANSWER 27 OF 82 EUROPATFULL COPYRIGHT 2004 WILA on STN  
TIEN Interfacially polymerized, bipiperidine-polyamide membranes for reverse  
osmosis and/or nanofiltration and process for making the same.
- L2 ANSWER 28 OF 82 EUROPATFULL COPYRIGHT 2004 WILA on STN  
TIEN PRODUCTION OF 1,2-DIHYDRO AND 2,2-DIHYDRO HEXAFLUOROPROPANES AND  
AZEOTROPES THEREOF WITH HF.
- L2 ANSWER 29 OF 82 EUROPATFULL COPYRIGHT 2004 WILA on STN  
TIEN Interfacially synthesized reverse osmosis membranes and processes for  
preparing the same.  
TIEN Interfacially synthesized reverse osmosis membranes and processes for  
preparing the same.
- L2 ANSWER 30 OF 82 USPATFULL on STN  
TI Catalysts for halogenated hydrocarbon processing, their precursors and  
their preparation and use
- L2 ANSWER 31 OF 82 USPATFULL on STN  
TI Process for the manufacture of 2-chloro-1,1,1-trifluoroethane
- L2 ANSWER 32 OF 82 PCTFULL COPYRIGHT 2004 Univentio on STN  
TIEN PROCESSES FOR THE PRODUCTION OF HEXAFLUOROPROPENE AND OPTIONAL OTHER  
HALOGENATED HYDROCARBONS CONTAINING FLUORINE  
TIFR PROCEDES RELATIFS A LA PRODUCTION D'HEXAFLUOROPROPENE ET EVENTUELLEMENT  
D'AUTRES HYDROCARBURES HALOGENES CONTENANT DU FLUOR
- L2 ANSWER 33 OF 82 PCTFULL COPYRIGHT 2004 Univentio on STN  
TIEN SELECTIVE MEMBRANE AND PROCESS FOR ITS PREPARATION  
TIFR MEMBRANE SELECTIVE ET PROCEDE DE PREPARATION DE CELLE-CI
- L2 ANSWER 34 OF 82 USPATFULL on STN  
TI Catalytic halogenated hydrocarbon processing and ruthenium  
catalysts for use therein
- L2 ANSWER 35 OF 82 USPATFULL on STN  
TI Process for the production of trifluoroethylene
- L2 ANSWER 36 OF 82 USPATFULL on STN  
TI Polymerization of, and depolymerization to, cyclic ethers using selected  
metal compound catalysts
- L2 ANSWER 37 OF 82 PCTFULL COPYRIGHT 2004 Univentio on STN

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TIEN CATALYSTS FOR HALOGENATED HYDROCARBON PROCESSING, THEIR PRECURSORS AND  
THEIR PREPARATION AND USE

TIFR CATALYSEURS DE TRAITEMENT D'HYDROCARBURES HALOGENES, LEURS PRECURSEURS,  
LEUR PREPARATION ET LEUR UTILISATION

L2 ANSWER 38 OF 82 PCTFULL COPYRIGHT 2004 Univentio on STN  
TIEN CATALYTIC HALOGENATED HYDROCARBON PROCESSING AND RUTHENIUM  
CATALYSTS FOR USE THEREIN

TIFR TRAITEMENT PAR CATALYSE DES HYDROCARBURES HALOGENES ET CATALYSEURS AU  
RUTHENIUM UTILISES

L2 ANSWER 39 OF 82 USPATFULL on STN  
TI Polymerization of, and depolymerization to, cyclic ethers using selected  
metal compound catalysts

L2 ANSWER 40 OF 82 EUROPATFULL COPYRIGHT 2004 WILA on STN  
TIEN PROCESS FOR THE MANUFACTURE OF 2-CHLORO-1,1,1-TRIFLUOROETHANE.

L2 ANSWER 41 OF 82 EUROPATFULL COPYRIGHT 2004 WILA on STN  
TIEN PROCESS FOR THE MANUFACTURE OF 2-CHLORO-1,1,1,2-TETRAFLUOROETHANE AND  
PENTAFLUOROETHANE.

L2 ANSWER 42 OF 82 EUROPATFULL COPYRIGHT 2004 WILA on STN  
TIEN PROCESS FOR THE MANUFACTURE OF 1,1,1,2-TETRAFLUOROETHANE.

L2 ANSWER 43 OF 82 EUROPATFULL COPYRIGHT 2004 WILA on STN  
TIEN PROCESS FOR THE MANUFACTURE OF 2,2-DICHLORO-1,1,1-TRIFLUOROETHANE,  
2-CHLORO-1,1,1,2-TETRAFLUOROETHANE AND PENTAFLUOROETHANE.

L2 ANSWER 44 OF 82 USPATFULL on STN  
TI Production of 1,2-dihydro and 2,2-dihydro hexafluoropropanes and  
azeotropes thereof with HF

L2 ANSWER 45 OF 82 USPATFULL on STN  
TI Acid gas fractionation process

L2 ANSWER 46 OF 82 USPATFULL on STN  
TI Acid gas fractionation process for fossil fuel gasifiers

L2 ANSWER 47 OF 82 USPATFULL on STN  
TI Process for manufacture of high purity 1, 1-dichlorotetrafluoroethane

L2 ANSWER 48 OF 82 USPATFULL on STN  
TI Polymerization of, and depolymerization to, cyclic ethers using selected  
metal compound catalysts

L2 ANSWER 49 OF 82 EUROPATFULL COPYRIGHT 2004 WILA on STN  
TIEN REGENERATION OR ACTIVATION OF NOBLE METAL CATALYSTS USING  
FLUOROHALOCARBONS OR FLUOROHALOHYDROCARBONS.

L2 ANSWER 50 OF 82 EUROPATFULL COPYRIGHT 2004 WILA on STN  
TIEN MANUFACTURE OF 1,1,1,2-TETRAFLUOROETHANE.

L2 ANSWER 51 OF 82 PCTFULL COPYRIGHT 2004 Univentio on STN  
TIEN PRODUCTION OF 1,2-DIHYDRO AND 2,2-DIHYDRO HEXAFLUOROPROPANES AND  
AZEOTROPES THEREOF WITH HF

TIFR PRODUCTION DE 1,2-DIHYDRO ET 2,2-DIHYDRO HEXAFLUOROPROPANES ET  
D'AZEOTROPES DE CES DERNIERS A L'AIDE DE FLUORURE D'HYDROGENE

L2 ANSWER 52 OF 82 PCTFULL COPYRIGHT 2004 Univentio on STN  
TIEN PROCESS FOR MANUFACTURE OF HIGH PURITY 1,1-DICHLOROTETRAFLUOROETHANE  
TIFR PROCEDE POUR PRODUIRE DU 1,1-DICHLOROTETRAFLUOROETHANE HAUTEMENT PUR

10/630,698

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TIEN MEMBRANE AND NON-MEMBRANE SOUR GAS TREATMENT PROCESS  
TIFR PROCEDE AVEC ET SANS MEMBRANE DE TRAITEMENT DE GAZ SULFUREUX
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TIFR PROCEDE DE TRAITEMENT DE GAZ SULFUREUX
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TIEN SOUR GAS MEMBRANE TREATMENT PROCESS INCLUDING DEHYDRATION  
TIFR PROCEDE DE TRAITEMENT MEMBRANAIRE DE GAZ SULFUREUX INCLUANT LA  
DESHYDRATATION
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TIEN POLYMERIZATION, AND DEPOLYMERIZATION, OF CYCLIC ETHERS USING  
HETEROGENEOUS CATALYSTS  
TIFR POLYMERISATION ET DEPOLYMERISATION D'ETHERS CYCLIQUES A L'AIDE DE  
CATALYSEURS HETEROGENES
- L2 ANSWER 57 OF 82 USPATFULL on STN  
TI Process for manufacture of high purity 1,1-dichlorotetrafluoroethane
- L2 ANSWER 58 OF 82 USPATFULL on STN  
TI Sour gas treatment process
- L2 ANSWER 59 OF 82 USPATFULL on STN  
TI Sour gas treatment process including membrane and non-membrane treatment steps
- L2 ANSWER 60 OF 82 USPATFULL on STN  
TI Sour gas treatment process including dehydration of the gas stream
- L2 ANSWER 61 OF 82 EUROPATFULL COPYRIGHT 2004 WILA on STN  
TIEN ACTIVATION OF NOBLE METAL CATALYSTS FOR USE IN HYDRODEHALOGENATION OF  
HALOGEN-SUBSTITUTED HYDROCARBONS CONTAINING FLUORINE AND AT LEAST ONE  
OTHER HALOGEN.
- L2 ANSWER 62 OF 82 USPATFULL on STN  
TI Manufacture of 1,1,1,2-tetrafluoroethane
- L2 ANSWER 63 OF 82 USPATFULL on STN  
TI Process for the manufacture of 1,1,1,2-tetrafluoroethane
- L2 ANSWER 64 OF 82 USPATFULL on STN  
TI Process for the manufacture of 2,2-dichloro-1,1,1-trifluoroethane,  
2-chloro-1,1,1,2-tetrafluoroethane and pentafluoroethane
- L2 ANSWER 65 OF 82 USPATFULL on STN  
TI Process for the manufacture of 2-chloro-1,1,1,2-tetrafluoroethane and  
pentafluoroethane
- L2 ANSWER 66 OF 82 EUROPATFULL COPYRIGHT 2004 WILA on STN  
TIEN Gas-phase fluorination process.  
TIEN Gas-phase fluorination process.
- L2 ANSWER 67 OF 82 USPATFULL on STN  
TI Interfacially synthesized reverse osmosis membranes and processes for  
preparing the same
- L2 ANSWER 68 OF 82 USPATFULL on STN  
TI Activation of noble metal catalysts for use in hydrodehalogenation of

10/630,698

halogen-substituted hydrocarbons containing fluorine and at least one other halogen

L2 ANSWER 69 OF 82 USPATFULL on STN  
TI Manufacture of 1,1,1,2-tetrafluoroethane

L2 ANSWER 70 OF 82 PCTFULL COPYRIGHT 2004 Univentio on STN  
TIEN PROCESS FOR REMOVING CONDENSABLE COMPONENTS FROM GAS STREAMS  
TIFR PROCEDE SERVANT A RETIRER DE FLUX GAZEUX DES CONSTITUANTS CONDENSABLES

L2 ANSWER 71 OF 82 PCTFULL COPYRIGHT 2004 Univentio on STN  
TIEN PROCESS FOR THE MANUFACTURE OF 2-CHLORO-1,1,1,2-TETRAFLUOROETHANE AND PENTAFLUOROETHANE  
TIFR PROCEDE DE FABRICATION DE 2-CHLORO-1,1,1,2-TETRAFLUOROETHANE ET PENTAFLUOROETHANE

L2 ANSWER 72 OF 82 PCTFULL COPYRIGHT 2004 Univentio on STN  
TIEN PROCESS FOR THE MANUFACTURE OF 1,1,1,2-TETRAFLUOROETHANE  
TIFR PROCEDE DE FABRICATION DE 1,1,1,2-TETRAFLUOROETHANE

L2 ANSWER 73 OF 82 PCTFULL COPYRIGHT 2004 Univentio on STN  
TIEN PROCESS FOR THE MANUFACTURE OF 2-CHLORO-1,1,1-TRIFLUOROETHANE  
TIFR PROCEDE DE FABRICATION DE 2-CHLORO-1,1,1-TRIFLUOROETHANE

L2 ANSWER 74 OF 82 PCTFULL COPYRIGHT 2004 Univentio on STN  
TIEN PROCESS FOR THE MANUFACTURE OF 2,2-DICHLORO-1,1,1-TRIFLUOROETHANE, 2-CHLORO-1,1,1,2-TETRAFLUOROETHANE AND PENTAFLUOROETHANE  
TIFR PROCEDE DE FABRICATION DE 2,2-DICHLORO-1,1,1-TRIFLUOROETHANE, 2-CHLORO-1,1,1,2-TETRAFLUOROETHANE ET PENTAFLUOROETHANE

L2 ANSWER 75 OF 82 USPATFULL on STN  
TI Activation of noble metal catalysts using fluorohalocarbons or fluorohalohydrocarbons

L2 ANSWER 76 OF 82 PCTFULL COPYRIGHT 2004 Univentio on STN  
TIEN ACTIVATION OF NOBLE METAL CATALYSTS FOR USE IN HYDRODEHALOGENATION OF HALOGEN-SUBSTITUTED HYDROCARBONS CONTAINING FLUORINE AND AT LEAST ONE OTHER HALOGEN  
TIFR ACTIVATION DE CATALYSEURS DE METAUX PRECIEUX DESTINES A L'HYDRODEHALOGENATION DES HYDROCARBURES SUBSTITUES PAR HALOGENE ET CONTENANT DU FLUOR ET AU MOINS UN AUTRE HALOGENE

L2 ANSWER 77 OF 82 PCTFULL COPYRIGHT 2004 Univentio on STN  
TIEN REGENERATION OR ACTIVATION OF NOBLE METAL CATALYSTS USING FLUOROHALOCARBONS OR FLUOROHALOHYDROCARBONS  
TIFR REGENERATION OU ACTIVATION D'UN CATALYSEUR EN METAL PRECIEUX A L'AIDE D'HALOCARBONES FLUORES OU D'HALOHYDROCARBONES FLUORES

L2 ANSWER 78 OF 82 USPATFULL on STN  
TI Regeneration of noble metal catalysts used in hydrodehalogenation of halogen-substituted hydrocarbons containing fluorine and at least one other halogen

L2 ANSWER 79 OF 82 PCTFULL COPYRIGHT 2004 Univentio on STN  
TIEN MANUFACTURE OF 1,1,1,2-TETRAFLUOROETHANE  
TIFR FABRICATION DE 1,1,1,2-TETRAFLUOROETHANE

L2 ANSWER 80 OF 82 USPATFULL on STN  
TI Regeneration or activation of noble metal catalysts using fluorohalocarbons or fluorohalohydrocarbons

L2 ANSWER 81 OF 82 USPATFULL on STN

10/630,698

TI Gas-phase fluorination process

L2 ANSWER 82 OF 82 JAPIO (C) 2004 JPO on STN  
TI METHOD FOR PRODUCING PERFLUOROPOLYETHERS HAVING ALDEHYDE, ALCOHOL, AND AMINE TERMINAL GROUPS

=> d 2,5,26,30,34,37,81,82 bib ab

L2 ANSWER 2 OF 82 CAPLUS COPYRIGHT 2004 ACS on STN  
AN 2004:117259 CAPLUS  
DN 140:146686  
TI Process for the preparation of perfluoropolyethers having aldehyde, alcohol, and amine end groups by catalytic reduction  
IN Di, Meo Antonello; Picozzi, Rosaldo; Tonelli, Claudio  
PA Solvay Solexis S.P.A., Italy  
SO Eur. Pat. Appl., 10 pp.  
CODEN: EPXXDW

DT Patent

LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	EP 1388556	A2	20040211	EP 2003-17183	20030729
	EP 1388556	A3	20040331		
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
	US 2004068144	A1	20040408	US 2003-630698	20030731
	JP 2004068007	A2	20040304	JP 2003-205414	20030801

PRAI IT 2002-MI1734 A 20020801

AB A process for the perfluoropolyether preparation having reactive end groups -CH<sub>2</sub>NH<sub>2</sub>, -CHO, -CH<sub>2</sub>OH, by reduction of the corresponding perfluoropolyethers having -CN, -COCl, -CHO end groups by using gaseous hydrogen in the presence of a catalyst constituted by Pd, Rh, or Ru, supported on solid metal fluorides, at 20-150° and under a pressure between 1 and 50 atmospheric is disclosed.

L2 ANSWER 5 OF 82 EUROPATFULL COPYRIGHT 2004 WILA on STN

GRANTED PATENT - ERTEILTES PATENT - BREVET DELIVRE

AN 1084093 EUROPATFULL ED 20040819 EW 200434 FS PS  
TIEN PROCESSES FOR THE PRODUCTION OF HEXAFLUOROPROPENE AND OPTIONALLY OTHER HALOGENATED HYDROCARBONS CONTAINING FLUORINE.  
TIDE VERFAHREN ZUR HERSTELLUNG VON HEXAFLUORPROPEN UND GEgebenenfalls WEITEREN HALOGENIERTEN FLUOR ENTHALTENDEN KOHLENWASSERSTOFFEN.  
TIFR PROCEDES RELATIFS A LA PRODUCTION D'HEXAFLUORPROPENE ET EVENTUELLEMENT D'AUTRES HYDROCARBURES HALOGENES CONTENANT DU FLUOR.  
IN SIEVERT, Allen, Capron, 215 Rhett Lane, Elkton, MD 21921, US;  
RAO, Velliyur, Nott, Mallikarjuna, 1 Georgetown Avenue, Wilmington, DE 19809, US;  
WALCZAK, Francis, J., 203 Jefferson Avenue, New Castle, DE 19720, US  
PA E.I. DUPONT DE NEMOURS AND COMPANY, 1007 Market Street, Wilmington, Delaware 19898, US  
PAN 2567250  
AG Towler, Philip Dean et al., Frank B. Dehn & Co., European Patent Attorneys, 179 Queen Victoria Street, London EC4V 4EL, GB  
AGN 75321  
OS MEPB2004035 EP 1084093 B1 0015  
SO Wila-EPS-2004-H34-T1  
DT Patent

10/630,698

LA Anmeldung in Englisch; Veröffentlichung in Englisch  
DS R DE; R FR; R GB; R IT; R NL  
PIT EPB1 EUROPÄISCHE PATENTSCHRIFT (Internationale Anmeldung)  
PI EP 1084093 B1 20040818  
OD 20010321  
AI EP 1999-928367 19990602  
PRAI US 1998-87751 19980602  
RLI WO 99-US12246 990602 INTAKZ  
WO 1999062851 991209 INTPNR  
REP EP 434407 A EP 434409 A  
WO 90-08748 A WO 97-19751 A  
GB 821211 A GB 2313118 A  
US 2576823 A US 5523501 A

L2 ANSWER 26 OF 82 USPATFULL on STN  
AN 2001:226805 USPATFULL  
TI Processes for the production of hexafluoropropene and optionally other  
halogenated hydrocarbons containing fluorine  
IN Sievert, Allen Capron, Elkton, MD, United States  
Rao, V. N. Mallikarjuna, Wilmington, DE, United States  
Walczak, Francis J., New Castle, DE, United States  
PA E. I. du Pont de Nemours and Company, Wilmington, DE, United States  
(U.S. corporation)  
PI US 6329559 B1 20011211  
WO 9962851 19991209  
AI US 2000-701448 20001127 (9)  
WO 1999-US12246 19990602  
20001127 PCT 371 date  
20001127 PCT 102(e) date  
PRAI US 1998-87751P 19980602 (60)  
PT Utility

PRAI US 1998-87751P 19980602 (60)

## DT Utility

FS GRANTED

EXNAM Primary Examiner: Siegel, Alan

CLMN Number of Claims: 20

ECL Exemplary Claim: 1

DRWN 1 Drawing Figure(s); 1 Drawing Page(s)

LN.CNT 961

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB A process is disclosed for the manufacture of CF.sub.3 CF.dbd.CF.sub.2 and optionally at least one compound selected from CF.sub.3 CH.sub.2 CF.sub.3 and CF.sub.3 CHFCHF.sub.2. The process involves contacting a reactor feed including a precursor stream of at least one halogenated propane of the formula CX.sub.3 CH.sub.2 CH.sub.y X.sub.(3-y) and/or halogenated propene of the formula CX.sub.3 CH.dbd.CH.sub.y X.sub.(2-y), where each X is Cl or F and y is 0, 1 or 2 (provided that the average fluorine content of the precursor stream is no more than 5 fluorine substituents per molecule) with HF and Cl.sub.2 in a chlorofluorination reaction zone containing a fluorination catalyst and operating at a temperature between about 150° C. and 400° C., to produce a reaction zone effluent including HF, HCl and a mixture of reaction products of the precursor feed which contains at least one compound of the formula C.sub.3 Cl.sub.2 F.sub.6 including CClF.sub.2 CClFCF.sub.3 and at least one compound of the formula C.sub.3 HC1F.sub.6, including CHF.sub.2 CClFCF.sub.3 and has an average fluorine content which is at least one fluorine substituent per molecule more than the average fluorine content of the precursor stream. The chlorofluorination reaction zone effluent is distilled to produce (i) a low-boiling component including HCl (and when they are present in the reaction zone effluent, C.sub.3 F.sub.8, C.sub.3 ClF.sub.7 and C.sub.3 HF.sub.7), (ii) a hydrogenation feed component containing at least one component of the formula C.sub.3 Cl.sub.2 F.sub.6 including CClF.sub.2 CClFCF.sub.3 and at least one compound of the formula C.sub.3

10/630,698

HClF.sub.6 including CHF.sub.2 CClFCF.sub.3, and an underfluorinated component including halogenated propanes containing at least one chlorine substituent and from one to five fluorine substituents. The CClF.sub.2 CClFCF.sub.3 and CHF.sub.2 CClFCF.sub.3 of hydrogenation feed component (ii) is reacted with hydrogen to produce a mixture including CF.sub.3 CF.dbd.CF.sub.2 and CF.sub.3 CHFCHF.sub.2 and the CF.sub.3 CF.dbd.CF.sub.2 from this product mixture is recovered. Underfluorinated component (iii) is returned to the chlorofluorination reaction zone.

L2 ANSWER 30 OF 82 USPATFULL on STN  
AN 2000:132057 USPATFULL  
TI Catalysts for halogenated hydrocarbon processing, their precursors and their preparation and use  
IN Duzick, Timothy C., Hockessin, DE, United States  
Rao, Velliyur Nott Mallikarjuna, Wilmington, DE, United States  
Subramanian, Munirpallam A., Kennett Square, PA, United States  
PA E. I. du Pont de Nemours and Company, Wilmington, DE, United States (U.S. corporation)  
PI US 6127585 20001003  
WO 9719751 19970605  
AI US 1998-77267 19980527 (9)  
WO 1996-US18967 19961126  
19980527 PCT 371 date  
19980527 PCT 102(e) date  
PRAI US 1995-7734P 19951129 (60)  
DT Utility  
FS Granted  
EXNAM Primary Examiner: Wu, David W.; Assistant Examiner: Zalukaeva, Tanya  
CLMN Number of Claims: 20  
ECL Exemplary Claim: 1  
DRWN No Drawings  
LN.CNT 958

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Processes are disclosed for decreasing the chlorine to carbon ratio for halogenated hydrocarbons containing chlorine and from 1 to 6 carbon atoms, in the presence of a multiphase catalyst. The processes each involve (1) preparing a single phase solid catalyst precursor which has a structure that collapses at a temperature of about 400° C. or less and has the formula (NH<sub>3</sub>)<sub>6</sub>Ru<sub>1-r-s</sub>Co<sub>r</sub>Cr<sub>s</sub>MF<sub>6</sub>, where r+s is in the range of 0.00 to 0.99, and M is at least one trivalent metal selected from the group consisting of Al, Cr, Fe, V, Sc and Ga; and (2) producing the multiphase catalyst by heating the single phase solid catalyst precursor to about 400° C. or less in an non-oxidizing atmosphere to produce a multiphase composition wherein a phase containing ruthenium is homogeneously dispersed with a phase containing metal fluoride.

Also disclosed are single phase fluoride compositions having the formula (NH<sub>3</sub>)<sub>6</sub>Ru<sub>1-r-s</sub>Co<sub>r</sub>Cr<sub>s</sub>MF<sub>6</sub>, where r+s is in the range of 0.00 to 0.99, and M is at least one trivalent element selected from the group consisting of Al, Cr, Fe, V, Sc and Ga; and multiphase catalyst compositions consisting essentially of metallic ruthenium and fluorides of at least one element selected from the group consisting of Al, Co, Cr, Fe, V, Sc and Ga, wherein the ruthenium is homogeneously dispersed with phases of the fluorides.

L2 ANSWER 34 OF 82 USPATFULL on STN  
AN 1999:75850 USPATFULL  
TI Catalytic halogenated hydrocarbon processing and ruthenium catalysts for use therein

10/630,698

IN Rao, Velliur Nott Mallikarjuna, Wilmington, DE, United States  
PA E. I. du Pont de Nemours and Company, Wilmington, DE, United States  
(U.S. corporation)

PI US 5919994 19990706  
WO 9719750 19970605

AI US 1997-875470 19970728 (8)  
WO 1996-US18952 19961126

19970728 PCT 371 date  
19970728 PCT 102(e) date

PRAI US 1995-7702P 19951129 (60)

DT Utility

FS Granted

EXNAM Primary Examiner: Yildirim, Bekir L.

CLMN Number of Claims: 8

ECL Exemplary Claim: 1

DRWN No Drawings

LN.CNT 1023

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Processes for decreasing the chlorine to carbon ratio for halogenated hydrocarbons containing chlorine and from 1 to 6 carbon atoms, in the presence of a catalyst are disclosed. The processes are each characterized by employing a catalyst comprising ruthenium on a support of (i) fluorided alumina, (ii) aluminum fluoride, or (iii) fluorides of Zn, Mg, Ca, Ba, Y, Sm, Eu, and/or Dy. Also disclosed are multiphase catalyst compositions of ruthenium supported on fluorides of Zn, Mg, Ca, Ba, Y, Sm, Eu and/or Dy.

L2 ANSWER 37 OF 82 PCTFULL COPYRIGHT 2004 Univentio on STN

AN 1997019751 PCTFULL ED 20020514

TIEN CATALYSTS FOR HALOGENATED HYDROCARBON PROCESSING, THEIR PRECURSORS AND THEIR PREPARATION AND USE

TIFR CATALYSEURS DE TRAITEMENT D'HYDROCARBURES HALOGENES, LEURS PRECURSEURS, LEUR PREPARATION ET LEUR UTILISATION

IN DUZICK, Timothy, C.;  
RAO, Velliur, Nott, Mallikarjuna;  
SUBRAMANIAN, Munirpallam, A.

PA E.I. DU PONT DE NEMOURS AND COMPANY;  
DUZICK, Timothy, C.;  
RAO, Velliur, Nott, Mallikarjuna;  
SUBRAMANIAN, Munirpallam, A.

LA English

DT Patent

PI WO 9719751 A1 19970605

DS W: JP US AT BE CH DE DK ES FI FR GB GR IE IT LU MC NL PT SE

AI WO 1996-US18967 A 19961126

PRAI US 1995-60/007,734 19951129

ABEN Processes are disclosed for decreasing the chlorine to carbon ratio for halogenated hydrocarbons containing chlorine and from 1 to 6 carbon atoms, in the presence of a multiphase catalyst. The processes each involve (1) preparing a single phase solid catalyst precursor which has a structure that collapses at a temperature of about 400 &deg;C or less and has the formula  $(NH_3)_6Ru^{r+s}Cl_xM^{(4-r-s)}F_6$ , where r+s is in the range of 0.00 to 0.99, and M is at least one trivalent metal selected from the group consisting of Al, Cr, Fe, V, Sc and Ga; and (2) producing the multiphase catalyst by heating the single phase solid catalyst precursor to about 400 &deg;C or less in a non-oxidizing atmosphere to produce a multiphase composition wherein a phase containing ruthenium

10/630,698

is homogeneously dispersed with a phase containing metal fluoride. Also disclosed are single phase fluoride compositions having the formula  $(\text{NH}_3)_6\text{Ru}^{1-r-s}\text{CorCrsMF}_6$ , where  $r+s$  is in the range of 0.00 to 0.99, and M is at least one trivalent element selected from the group consisting of Al, Cr, Fe, V, Sc and Ga; and multiphase catalyst compositions consisting essentially of metallic ruthenium and fluorides of at least one element selected from the group consisting of Al, Co, Cr, Fe, V, Sc and Ga, wherein the ruthenium is homogeneously dispersed with phases of the fluorides.

ABFR L'invention concerne des procedes servant a diminuer le rapport entre le chlore et le carbone pour des hydrocarbures halogenes contenant du chlore et de 1 a 6 atomes de carbone, en presence d'un catalyseur a phases multiples. Ces procedes consistent chacun (1) a preparer un precurseur de catalyseur solide monophase, dont la structure s'affaisse a une temperature egale ou inferieure a 400 &deg;C et qui possede la formule  $(\text{NH}_3)_6\text{Ru}^{1-r-s}\text{CorCrsMF}_6$  dans laquelle  $r+s$  est situe dans la plage de 0,00 a 0,99 et M represente au moins un metal trivalent selectionne dans le groupe constitue par Al, Cr, Fe, V, Sc et Ga et (2) a produire le catalyseur a phases multiples par rechauffement du catalyseur solide monophase a une temperature egale ou inferieure a 400 &deg;C dans une atmosphere non oxydante, afin d'obtenir une composition a phases multiples dans laquelle une phase contenant ruthenium est dispersee de facon homogene avec une phase contenant fluorure metallique. L'invention concerne également des compositions monophases de fluor possedant la formule  $(\text{NH}_3)_6\text{RU}^{1-r-s}\text{CorCrsMF}_6$  dans laquelle  $r+s$  est situe dans la plage de 0,00 a 0,99 et M represente au moins un element trivalent selectionne dans le groupe constitue par Al, Cr, Fe, V, Sc et Ga, ainsi que des compositions de catalyseur a phases multiples constituees essentiellement par ruthenium et des fluorures metalliques d'au moins un element selectionne dans le groupe constitue par Al, Co, Cr, Fe, V, Sc et Ga, le ruthenium etant disperse de facon homogene avec des phases des fluorures.

L2 ANSWER 81 OF 82 USPATFULL on STN  
AN 90:34289 USPATFULL  
TI Gas-phase fluorination process  
IN Manzer, Leo E., Wilmington, DE, United States  
PA E. I. du Pont de Nemours and Company, Wilmington, DE, United States  
(U.S. corporation)  
PI US 4922037 19900501  
AI US 1989-355867 19890519 (7)  
RLI Continuation of Ser. No. US 1988-160003, filed on 24 Feb 1988, now abandoned  
DT Utility  
FS Granted  
EXNAM Primary Examiner: Evans, J. E.  
LREP Shipley, James E.  
CLMN Number of Claims: 13  
ECL Exemplary Claim: 1  
DRWN No Drawings

10/630,698

LN.CNT 397

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB An improved process for the manufacture of 1,1,1,2-tetrafluoroethane, more particularly, a gas-phase reaction of a 1,1,1-trifluorochloroethane with hydrogen fluoride in the presence of a selected metal on aluminum fluoride or carbon.

L2 ANSWER 82 OF 82 JAPIO (C) 2004 JPO on STN

AN 2004-068007 JAPIO

TI METHOD FOR PRODUCING PERFLUOROPOLYETHERS HAVING ALDEHYDE, ALCOHOL, AND AMINE TERMINAL GROUPS

IN DI MEO ANTONELLO; PICOZZI ROSALDO; TONELLI CLAUDIO

PA SOLVAY SOLEXIS SPA

PI JP 2004068007 A 20040304 Heisei

AI JP 2003-205414 (JP2003205414 Heisei) 20030801

PRAI IT 2002-MI02 1734 20020801

SO PATENT ABSTRACTS OF JAPAN (CD-ROM), Unexamined Applications, Vol. 2004

AB PROBLEM TO BE SOLVED: To provide a method for producing a reduced compound having a corresponding aldehyde, alcohol, or amine terminal group in a high yield of >=90% from precursors of perfluoropolyethers having an acyl-chloride, aldehyde or nitrile terminal group.

SOLUTION: The problem is solved by the method for producing a perfluoropolyether having a reactive terminal group of

-CH<SB>2</SB>NH<SB>2</SB>, -CHO, or CH<SB>2</SB>OH by reducing

the corresponding perfluoropolyether having a -CN, -COCl, or

-CHO terminal group using a hydrogen gas in the

presence of a catalyst constituted with Pd, Rh, and

Ru supported on a solid metallic fluoride at a

temperature of 20-150&deg;C under a pressure of 1-50 atmospheric

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status data from INPADOC  
NEWS 9 SEP 01 INPADOC: New family current-awareness alert (SDI) available  
NEWS 10 SEP 01 New pricing for the Save Answers for SciFinder Wizard within  
STN Express with Discover!  
NEWS 11 SEP 01 New display format, HITSTR, available in WPIDS/WPINDEX/WPIX  
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NEWS 13 SEP 27 SWETSCAN will no longer be available on STN  
NEWS 14 OCT 28 KOREAPAT now available on STN

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DICTIONARY FILE UPDATES: 8 NOV 2004 HIGHEST RN 777024-10-9

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P<sup>1</sup>  
2CF2CF3 5CH<sub>2</sub>OH 1<sup>1</sup>  
3CF2CF3 6CH<sub>2</sub>NH<sub>2</sub> 2<sub>2</sub> 7 5<sup>3</sup> 10  
Cl<sup>4</sup><sub>3</sub> CH<sub>3</sub> 3<sub>3</sub> 4 6 6<sup>9</sup> 11  
5 4 12<sup>7</sup>

chain nodes :  
1 2 3 4 5 6 7 8 9 10 11 12 21 22 23 25 26 27 28 29 30 31 32  
33 34 35 36 37 38 39 40 41 42 43 45  
chain bonds :  
2-7 3-4 4-6 8-10 9-11 21-22 22-23 22-25 22-26 26-27 27-28 28-29 29-30  
30-31 31-32 31-33 31-34 34-35 35-36 36-37 37-40 37-38 37-39 40-41 41-42  
41-43 41-45

10/630,698

exact/norm bonds :  
21-22 22-25 22-26 31-34 36-37 37-40 40-41 41-43 41-45  
exact bonds :  
2-7 3-4 4-6 8-10 9-11 22-23 26-27 27-28 28-29 29-30 30-31 31-32 31-33  
34-35 35-36 37-38 37-39 41-42

G1:[\*1], [\*2], [\*3], [\*4], [\*5], [\*6], [\*7]

G2:[\*1], [\*2]

Match level :  
1:CLASS 2:CLASS 3:CLASS 4:CLASS 5:CLASS 6:CLASS 7:CLASS 8:CLASS 9:CLASS  
10:CLASS 11:CLASS 12:CLASS 21:CLASS 22:CLASS 23:CLASS 25:CLASS 26:CLASS  
27:CLASS 28:CLASS 29:CLASS 30:CLASS 31:CLASS 32:CLASS 33:CLASS 34:CLASS  
35:CLASS 36:CLASS 37:CLASS 38:CLASS 39:CLASS 40:CLASS 41:CLASS 42:CLASS  
43:CLASS 45:CLASS

L2 STRUCTURE UPLOADED

=> que L2 NOT L1

L3 QUE L2 NOT L1

=> s 13

STRUCTURE TOO LARGE - SEARCH ENDED

A structure in your query is too large. You may delete attributes or atoms to reduce the size of the structure and try again.

=> file stnguide

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	0.84	1.26

FILE 'STNGUIDE' ENTERED AT 06:58:40 ON 10 NOV 2004

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FILE CONTAINS CURRENT INFORMATION.

LAST RELOADED: Nov 5, 2004 (20041105/UP).

=> file reg

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	0.36	1.62

FILE 'REGISTRY' ENTERED AT 07:02:23 ON 10 NOV 2004

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STRUCTURE FILE UPDATES: 8 NOV 2004 HIGHEST RN 777024-10-9  
DICTIONARY FILE UPDATES: 8 NOV 2004 HIGHEST RN 777024-10-9

10/630,698

TSCA INFORMATION NOW CURRENT THROUGH MAY 21, 2004

Please note that search-term pricing does apply when conducting SmartSELECT searches.

Crossover limits have been increased. See HELP CROSSOVER for details.

Experimental and calculated property data are now available. For more information enter HELP PROP at an arrow prompt in the file or refer to the file summary sheet on the web at:  
<http://www.cas.org/ONLINE/DBSS/registryss.html>

=> ....Testing the current file.... screen

ENTER SCREEN EXPRESSION OR (END) :end

=> screen 1994 OR 2016 OR 2021 OR 2026 OR 1929 OR 1838

L4 SCREEN CREATED

=>  
Uploading C:\Program Files\Stnexp\Queries\10630698a.str

G2  
G1 O CFCF2O1-5CF2 F CF3 O0-5CFO1-5 F G2 25  
F F F F G12J226272829 30 33 3134 3536 39 38 40 41 43 45  
23 32 32 38 42



chain nodes :

1 2 3 4 5 6 7 8 9 10 11 12 21 22 23 25 26 27 28 29 30 31 32  
33 34 35 36 37 38 39 40 41 42 43 45

chain bonds :

2-7 3-4 4-6 8-10 9-11 21-22 22-23 22-25 22-26 26-27 27-28 28-29 29-30  
30-31 31-32 31-33 31-34 34-35 35-36 36-37 37-40 37-38 37-39 40-41 41-42  
41-43 41-45

exact/norm bonds :

21-22 22-25 22-26 31-34 36-37 37-40 40-41 41-43 41-45

exact bonds :

2-7 3-4 4-6 8-10 9-11 22-23 26-27 27-28 28-29 29-30 30-31 31-32 31-33  
34-35 35-36 37-38 37-39 41-42

10/630,698

G1:[\*1], [\*2], [\*3], [\*4], [\*5], [\*6], [\*7]

G2:[\*1], [\*2]

Match level :

1:CLASS 2:CLASS 3:CLASS 4:CLASS 5:CLASS 6:CLASS 7:CLASS 8:CLASS 9:CLASS  
10:CLASS 11:CLASS 12:CLASS 21:CLASS 22:CLASS 23:CLASS 25:CLASS 26:CLASS  
27:CLASS 28:CLASS 29:CLASS 30:CLASS 31:CLASS 32:CLASS 33:CLASS 34:CLASS  
35:CLASS 36:CLASS 37:CLASS 38:CLASS 39:CLASS 40:CLASS 41:CLASS 42:CLASS  
43:CLASS 45:CLASS

L5 STRUCTURE UPLOADED

=> que L5 NOT L4

L6 QUE L5 NOT L4

=> s 16

STRUCTURE TOO LARGE - SEARCH ENDED

A structure in your query is too large. You may delete attributes or atoms to reduce the size of the structure and try again.

=> file stnguide

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	0.84	2.46

FILE 'STNGUIDE' ENTERED AT 07:03:35 ON 10 NOV 2004

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FILE CONTAINS CURRENT INFORMATION.

LAST RELOADED: Nov 5, 2004 (20041105/UP).

=> file reg

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	0.30	2.76

FILE 'REGISTRY' ENTERED AT 07:06:51 ON 10 NOV 2004

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STRUCTURE FILE UPDATES: 8 NOV 2004 HIGHEST RN 777024-10-9  
DICTIONARY FILE UPDATES: 8 NOV 2004 HIGHEST RN 777024-10-9

TSCA INFORMATION NOW CURRENT THROUGH MAY 21, 2004

Please note that search-term pricing does apply when conducting SmartSELECT searches.

Crossover limits have been increased. See HELP CROSSOVER for details.

10/630,698

Experimental and calculated property data are now available. For more information enter HELP PROP at an arrow prompt in the file or refer to the file summary sheet on the web at:  
<http://www.cas.org/ONLINE/DBSS/registryss.html>

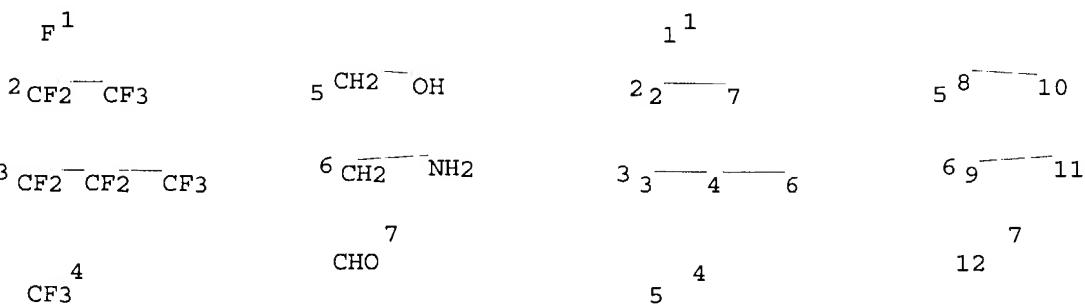
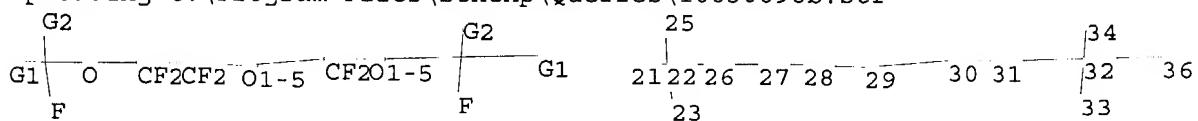
=> ....Testing the current file.... screen

ENTER SCREEN EXPRESSION OR (END) :end

=> screen 1994 OR 2016 OR 2021 OR 2026 OR 1929 OR 1838

L7 SCREEN CREATED

=>  
Uploading C:\Program Files\Stnexp\Queries\10630698b.str



chain nodes :  
1 2 3 4 5 6 7 8 9 10 11 12 21 22 23 25 26 27 28 29 30 31 32  
33 34 36

chain bonds :

2-7 3-4 4-6 8-10 9-11 21-22 22-23 22-25 22-26 26-27 27-28 28-29 29-30  
30-31 31-32 32-36 32-33 32-34

exact/norm bonds :

21-22 22-25 22-26 31-32 32-36 32-34

exact bonds :

2-7 3-4 4-6 8-10 9-11 22-23 26-27 27-28 28-29 29-30 30-31 32-33

10/630,698

G1:[\*1], [\*2], [\*3], [\*4], [\*5], [\*6], [\*7]

G2:[\*1], [\*2]

Match level :

1:CLASS 2:CLASS 3:CLASS 4:CLASS 5:CLASS 6:CLASS 7:CLASS 8:CLASS 9:CLASS  
10:CLASS 11:CLASS 12:CLASS 21:CLASS 22:CLASS 23:CLASS 25:CLASS 26:CLASS  
27:CLASS 28:CLASS 29:CLASS 30:CLASS 31:CLASS 32:CLASS 33:CLASS 34:CLASS  
36:CLASS

L8 STRUCTURE uploaded

=> que L8 NOT L7

L9 QUE L8 NOT L7

=> d

L9 HAS NO ANSWERS

L7 SCR 1994 OR 2016 OR 2021 OR 2026 OR 1929 OR 1838  
L8 STR

\* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT \*

Structure attributes must be viewed using STN Express query preparation.

L9 QUE L8 NOT L7

=> s 19

SAMPLE SEARCH INITIATED 07:07:31 FILE 'REGISTRY'  
SAMPLE SCREEN SEARCH COMPLETED - 0 TO ITERATE

100.0% PROCESSED 0 ITERATIONS 0 ANSWERS  
SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE \*\*COMPLETE\*\*  
BATCH \*\*COMPLETE\*\*

PROJECTED ITERATIONS: 0 TO 0

PROJECTED ANSWERS: 0 TO 0

L10 0 SEA SSS SAM L8 NOT L7

=> s 19 ful  
FULL SEARCH INITIATED 07:07:40 FILE 'REGISTRY'  
FULL SCREEN SEARCH COMPLETED - 0 TO ITERATE

100.0% PROCESSED 0 ITERATIONS 0 ANSWERS  
SEARCH TIME: 00.00.01

L11 0 SEA SSS FUL L8 NOT L7

=> file stnguide  
COST IN U.S. DOLLARS SINCE FILE TOTAL  
FULL ESTIMATED COST ENTRY SESSION  
156.26 159.02

FILE 'STNGUIDE' ENTERED AT 07:08:21 ON 10 NOV 2004  
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10/630,698

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FILE CONTAINS CURRENT INFORMATION.  
LAST RELOADED: Nov 5, 2004 (20041105/UP).

=> file reg		SINCE FILE ENTRY	TOTAL SESSION
COST IN U.S. DOLLARS			
FULL ESTIMATED COST		0.54	159.56

FILE 'REGISTRY' ENTERED AT 07:13:48 ON 10 NOV 2004  
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STRUCTURE FILE UPDATES: 8 NOV 2004 HIGHEST RN 777024-10-9  
DICTIONARY FILE UPDATES: 8 NOV 2004 HIGHEST RN 777024-10-9

TSCA INFORMATION NOW CURRENT THROUGH MAY 21, 2004

Please note that search-term pricing does apply when  
conducting SmartSELECT searches.

Crossover limits have been increased. See HELP CROSSOVER for details.

Experimental and calculated property data are now available. For more  
information enter HELP PROP at an arrow prompt in the file or refer  
to the file summary sheet on the web at:  
<http://www.cas.org/ONLINE/DBSS/registryss.html>

=> ....Testing the current file.... screen

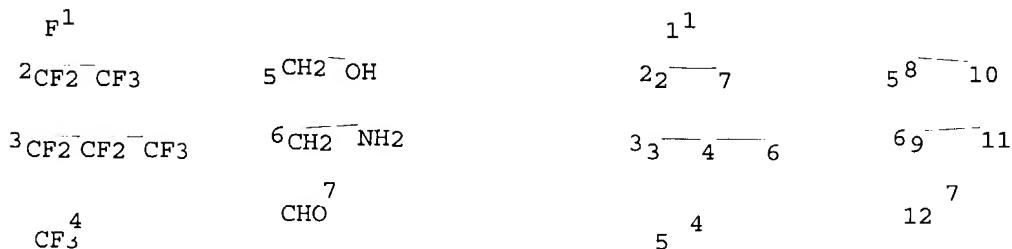
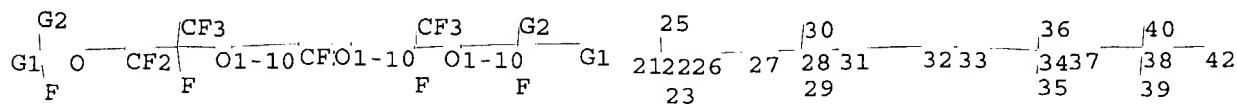
ENTER SCREEN EXPRESSION OR (END) :end

=> screen 1994 OR 2016 OR 2021 OR 2026 OR 1929 OR 1838

L12 SCREEN CREATED

=>  
Uploading C:\Program Files\Stnexp\Queries\10630698c.str

10/630,698



chain nodes :

1 2 3 4 5 6 7 8 9 10 11 12 21 22 23 25 26 27 28 29 30 31 32  
33 34 35 36 37 38 39 40 42

chain bonds :

2-7 3-4 4-6 8-10 9-11 21-22 22-23 22-25 22-26 26-27 27-28 28-29 28-30  
28-31 31-32 32-33 33-34 34-37 34-35 34-36 37-38 38-39 38-40 38-42

exact/norm bonds :

21-22 22-25 22-26 28-31 33-34 34-37 37-38 38-40 38-42

exact bonds :

2-7 3-4 4-6 8-10 9-11 22-23 26-27 27-28 28-29 28-30 31-32 32-33 34-35  
34-36 38-39

G1: [\*1], [\*2], [\*3], [\*4], [\*5], [\*6], [\*7]

G2: [\*1], [\*2]

Match level :

1:CLASS 2:CLASS 3:CLASS 4:CLASS 5:CLASS 6:CLASS 7:CLASS 8:CLASS 9:CLASS  
10:CLASS 11:CLASS 12:CLASS 21:CLASS 22:CLASS 23:CLASS 25:CLASS 26:CLASS  
27:CLASS 28:CLASS 29:CLASS 30:CLASS 31:CLASS 32:CLASS 33:CLASS 34:CLASS  
35:CLASS 36:CLASS 37:CLASS 38:CLASS 39:CLASS 40:CLASS 42:CLASS

L13 STRUCTURE UPLOADED

=> que L13 NOT L12

L14 QUE L13 NOT L12

10/630, 698

=> S 114  
STRUCTURE TOO LARGE - SEARCH ENDED

A structure in your query is too large. You may delete attributes or atoms to reduce the size of the structure and try again.

=> file stnguide  
COST IN U.S. DOLLARS  
  
FULL ESTIMATED COST

SINCE FILE ENTRY	TOTAL SESSION 0.84 160.40
---------------------	---------------------------------

FILE 'STNGUIDE' ENTERED AT 07:14:45 ON 10 NOV 2004  
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FILE CONTAINS CURRENT INFORMATION.  
LAST RELOADED: Nov 5, 2004 (20041105/UP).

=> file reg  
COST IN U.S. DOLLARS  
  
FULL ESTIMATED COST

SINCE FILE ENTRY	TOTAL SESSION 0.30
---------------------	--------------------------

FILE 'REGISTRY' ENTERED AT 07:17:49 ON 10 NOV 2004  
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STRUCTURE FILE UPDATES: 8 NOV 2004 HIGHEST RN 777024-10-9  
DICTIONARY FILE UPDATES: 8 NOV 2004 HIGHEST RN 777024-10-9

TSCA INFORMATION NOW CURRENT THROUGH MAY 21, 2004

Please note that search-term pricing does apply when conducting SmartSELECT searches.

Crossover limits have been increased. See HELP CROSSOVER for details.

Experimental and calculated property data are now available. For more information enter HELP PROP at an arrow prompt in the file or refer to the file summary sheet on the web at:  
<http://www.cas.org/ONLINE/DBSS/registryss.html>

=> ....Testing the current file.... screen

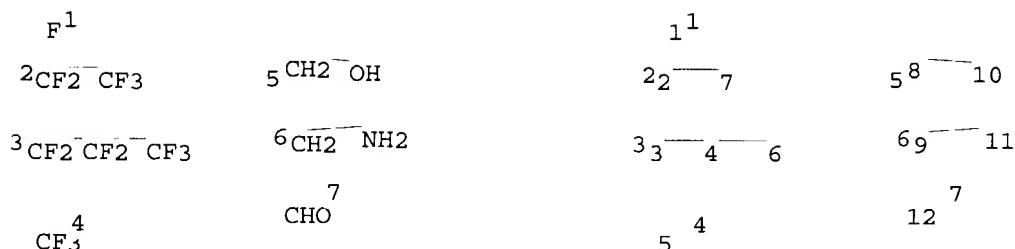
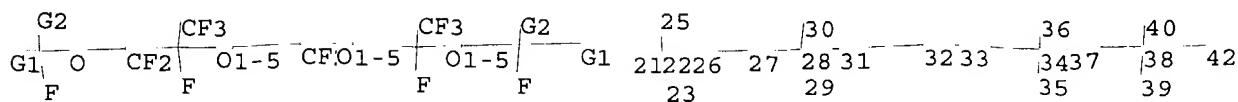
ENTER SCREEN EXPRESSION OR (END) :end

=> screen 1994 OR 2016 OR 2021 OR 2026 OR 1838

L15 SCREEN CREATED

=>  
Uploading C:\Program Files\Stnexp\Queries\10630698d.str

10/630,698



chain nodes :

1 2 3 4 5 6 7 8 9 10 11 12 21 22 23 25 26 27 28 29 30 31 32  
33 34 35 36 37 38 39 40 42

chain bonds :

2-7 3-4 4-6 8-10 9-11 21-22 22-23 22-25 22-26 26-27 27-28 28-29 28-30  
28-31 31-32 32-33 33-34 34-37 34-35 34-36 37-38 38-39 38-40 38-42

exact/norm bonds :

21-22 22-25 22-26 28-31 33-34 34-37 37-38 38-40 38-42

exact bonds :

2-7 3-4 4-6 8-10 9-11 22-23 26-27 27-28 28-29 28-30 31-32 32-33 34-35  
34-36 38-39

G1: [\*1], [\*2], [\*3], [\*4], [\*5], [\*6], [\*7]

G2: [\*1], [\*2]

Match level :

1:CLASS 2:CLASS 3:CLASS 4:CLASS 5:CLASS 6:CLASS 7:CLASS 8:CLASS 9:CLASS  
10:CLASS 11:CLASS 12:CLASS 21:CLASS 22:CLASS 23:CLASS 25:CLASS 26:CLASS  
27:CLASS 28:CLASS 29:CLASS 30:CLASS 31:CLASS 32:CLASS 33:CLASS 34:CLASS  
35:CLASS 36:CLASS 37:CLASS 38:CLASS 39:CLASS 40:CLASS 42:CLASS

L16 STRUCTURE UPLOADED

=> que L16 NOT L15

L17 QUE L16 NOT L15

10/630,698

=> s 117

SAMPLE SEARCH INITIATED 07:18:20 FILE 'REGISTRY'  
SAMPLE SCREEN SEARCH COMPLETED - 174 TO ITERATE

100.0% PROCESSED 174 ITERATIONS ( 10 INCOMPLETE) 10 ANSWERS  
SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE \*\*COMPLETE\*\*  
BATCH \*\*COMPLETE\*\*

PROJECTED ITERATIONS: 2689 TO 4271  
PROJECTED ANSWERS: 11 TO 389

L18 10 SEA SSS SAM L16 NOT L15

=> s 117 ful

FULL SEARCH INITIATED 07:18:29 FILE 'REGISTRY'  
FULL SCREEN SEARCH COMPLETED - 3484 TO ITERATE

100.0% PROCESSED 3484 ITERATIONS ( 156 INCOMPLETE) 156 ANSWERS  
SEARCH TIME: 00.00.07

L19 156 SEA SSS FUL L16 NOT L15

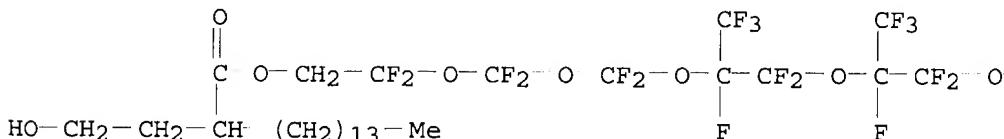
=> d scan

L19 156 ANSWERS REGISTRY COPYRIGHT 2004 ACS on STN  
ITERATION INCOMPLETE

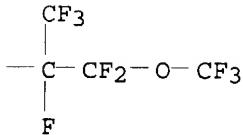
IN Hexadecanoic acid, 2-(2-hydroxyethyl)-, 2,2,4,4,6,6,8,9,9,11,12,12,14,15,1  
5,17,17,17-octadecafluoro-8,11,14-tris(trifluoromethyl)-3,5,7,10,13,16-  
hexaoxaheptadec-1-yl ester (9CI)

MF C32 H37 F27 O9

PAGE 1-A



PAGE 1-B



\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):file caplus casreact uspatful  
'FILE CAPLUS CASREACT USPATFUL' IS NOT VALID HERE

To display more answers, enter the number of answers you would like to see. To end the display, enter "NONE", "N", "0", or "END".  
HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):0

10/630,698

=> file caplus casreact uspatful

COST IN U.S. DOLLARS

SINCE FILE

ENTRY

TOTAL

FULL ESTIMATED COST

156.68

SESSION

317.38

FILE 'CAPLUS' ENTERED AT 07:20:01 ON 10 NOV 2004

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FILE 'CASREACT' ENTERED AT 07:20:01 ON 10 NOV 2004

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FILE 'USPATFULL' ENTERED AT 07:20:01 ON 10 NOV 2004

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=> s 119

L20 120 L19

=> dup rem 120

PROCESSING COMPLETED FOR L20

L21 108 DUP REM L20 (12 DUPLICATES REMOVED)

=> d 1-108 ti

L21 ANSWER 1 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Manufacture of magnetic recording media

L21 ANSWER 2 OF 108 USPATFULL on STN

TI Novel aerosol formulation containing a polar fluorinated molecule

L21 ANSWER 3 OF 108 USPATFULL on STN

TI Amphiphilic lubricants for magnetic media

L21 ANSWER 4 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN DUPLICATE 1

TI Amphiphilic lubricants with central polar polyfunctional group and pair of fluoroalkylether endgroups as topcoats for magnetic recording media

L21 ANSWER 5 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN

TI Aerosol formulation containing a polar fluorinated compound

L21 ANSWER 6 OF 108 USPATFULL on STN

TI Process for thermal decomposition of hexafluoropropylene oxide oligomers

L21 ANSWER 7 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN DUPLICATE 2

TI Perfluorinated organo substituted cyclosiloxanes and copolymers prepared from these cyclosiloxanes

L21 ANSWER 8 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN DUPLICATE 3

TI Perfluorinated ether organo substituted cyclosiloxanes and siloxane (co)polymers prepared from these cyclosiloxanes

L21 ANSWER 9 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN

TI Magnetic recording medium with fluorine-containing alkylcarboxylic acid lubricating layer

L21 ANSWER 10 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN

TI Process for thermal decomposition of hexafluoropropylene oxide oligomers

L21 ANSWER 11 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN

10/630,698

- TI Process for preparation of hexafluoropropylene oxide by oxidation of hexafluoropropylene
- L21 ANSWER 12 OF 108 USPATFULL on STN  
TI Fluoroalkylated amphiphilic ligands, their metallic complexes and their uses
- L21 ANSWER 13 OF 108 USPATFULL on STN  
TI Amides and esters of perfluoropolyoxaalkylene-sulfo- or perfluoropolyoxaalkylene-carboxylic acids and a process for producing same
- L21 ANSWER 14 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Water Core within Perfluoropolyether-Based Microemulsions Formed in Supercritical Carbon Dioxide
- L21 ANSWER 15 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Adsorption of fluorine-containing surfactants from aqueous solutions on the surface of polyamide fibers
- L21 ANSWER 16 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Radical additions to fluoroolefins. Photochemical mono-fluoroalkylation and sequential bis-fluoroalkylation of oxolane
- L21 ANSWER 17 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Effect of colloidal-chemical properties of fluorine-containing latexes and fluorocarbon surfactants on the modification of textiles
- L21 ANSWER 18 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN DUPLICATE 4  
TI Fluorine-containing alkylsuccinic acid diester and its preparation and use as a lubricant for magnetic recording media
- L21 ANSWER 19 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Manufacture of amides and esters of perfluoropolyoxyalkylenesulfonic or -carboxylic acids
- L21 ANSWER 20 OF 108 USPATFULL on STN  
TI Fluoroalkylated amphiphilic ligands and their metallic complexes
- L21 ANSWER 21 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Reaction of perfluoropolyoxapolypropenecarboxylic acids with metal carbonates and acid fluorides with 3-Amino-1,2,4-Triazole
- L21 ANSWER 22 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Esterification of perfluoropolyoxapolypropylenecarboxylic acid (n = 8)
- L21 ANSWER 23 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Fluoroalkylated amphiphilic ligands, their metallic complexes and their uses
- L21 ANSWER 24 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Preparation of fluoroalkanoic acid esters and magnetic recording medium with lubricant layer containing them
- L21 ANSWER 25 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Preparation of fluorinated alcohols and magnetic recording media using them as lubricants
- L21 ANSWER 26 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Perfluorooxalkyl group-containing polymers
- L21 ANSWER 27 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN

10/630,698

- TI Perfluorooxalkyl group-terminated vinyl polymers
- L21 ANSWER 28 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Preparation of polyfluoroalkanoyl peroxides as polymerization initiators
- L21 ANSWER 29 OF 108 USPATFULL on STN  
TI Polyfluoroalkanoyl peroxide
- L21 ANSWER 30 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Surface activity of fluorine-containing surfactants in polar solvents and water-organic mixtures
- L21 ANSWER 31 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Preparation of perfluoroacetal and perfluoroketal compounds and use thereof in thermal shock testing
- L21 ANSWER 32 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Methylcarbinol-terminated hexafluoropropylene oxide oligoether derivatives
- L21 ANSWER 33 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Preparation of carbonyl fluorides by oligomerization of hexafluoropropene oxides
- L21 ANSWER 34 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Process and catalysts for the manufacture of hexafluoropropylene oxide oligomers
- L21 ANSWER 35 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Electrolytic decarboxylation of perfluorocarboxylic acids or their soluble salts and subsequent dimerization of the radicals produced
- L21 ANSWER 36 OF 108 USPATFULL on STN  
TI Process for the oligomerization of hexafluoropropene oxide
- L21 ANSWER 37 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Aggregation of perfluorinated polymers in aqueous solution studied by ESR
- L21 ANSWER 38 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Process for preparation of perfluorinated carboxylic acid fluorides
- L21 ANSWER 39 OF 108 USPATFULL on STN  
TI Process for the preparation of perfluorinated carbonyl fluorides
- L21 ANSWER 40 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN DUPLICATE 5  
TI Preparation of carbonyl fluoride compounds
- L21 ANSWER 41 OF 108 USPATFULL on STN  
TI Fluoropolyethers containing end groups endowed with anchoring capacity
- L21 ANSWER 42 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Fuel cells
- L21 ANSWER 43 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN DUPLICATE 6  
TI Fluorine-containing methacrylate esters
- L21 ANSWER 44 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Fluorine-containing polymers with oxygen permeability for medical use
- L21 ANSWER 45 OF 108 USPATFULL on STN  
TI Shaped article of synthetic resin having improved surface
- L21 ANSWER 46 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN

10/630,698

- TI Synthetic resin films with water and oil repellence
- L21 ANSWER 47 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Lubricant finishes
- L21 ANSWER 48 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Fluoropolyether compounds
- L21 ANSWER 49 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Surface active substances containing an oligo(hexafluoropropene oxide) chain as a hydrophobic oleophobic moiety
- L21 ANSWER 50 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Acrylic acid esters
- L21 ANSWER 51 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Water and oil repellents
- L21 ANSWER 52 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Synthesis of fluorinated surfactants containing hexafluoropropene oxide as a hydrophobic group and properties of the solutions
- L21 ANSWER 53 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Polymerization of fluorine-containing monomers
- L21 ANSWER 54 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Foaming agent for extinguishing fires
- L21 ANSWER 55 OF 108 USPATFULL on STN  
TI Alkyl perfluoro- $\omega$ -fluoroformyl esters and their preparation
- L21 ANSWER 56 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Methods of calculating engineering parameters for gas separations
- L21 ANSWER 57 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN DUPLICATE 7  
TI Alkyl perfluoro- $\omega$ -fluoroformyl esters and monomers therefrom
- L21 ANSWER 58 OF 108 USPATFULL on STN  
TI Process for the preparation of fluorine-containing ketones
- L21 ANSWER 59 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Alkylperfluoro- $\omega$ -fluoroformyl esters
- L21 ANSWER 60 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Perfluoro ketones
- L21 ANSWER 61 OF 108 USPATFULL on STN  
TI Alkyl perfluoro- $\omega$ -fluoroformyl esters and their preparation
- L21 ANSWER 62 OF 108 USPATFULL on STN  
TI Fluorocarbon triazine polymers
- L21 ANSWER 63 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Synthesis of perfluoro(polyether) difunctional compounds
- L21 ANSWER 64 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Methods for the estimation of vapor pressures and oxygen solubilities of fluoroochemicals for possible application in artificial blood formulations
- L21 ANSWER 65 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN DUPLICATE 8  
TI Fluorocarbon dye dispersion for exhaust disperse dyeing

10/630,698

- L21 ANSWER 66 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Fluorocarbon triazine polymers
- L21 ANSWER 67 OF 108 USPATFULL on STN  
TI Fluoroalkyleneether difunctional compounds
- L21 ANSWER 68 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Study of the kinetics of the reaction of hexafluoropropylene oxide with organic salts in a medium of aprotic solvents
- L21 ANSWER 69 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Oligomeric fluorinated additives as surface modifiers for solid polymers
- L21 ANSWER 70 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI The solubility of oxygen in highly fluorinated liquids
- L21 ANSWER 71 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN DUPLICATE 9  
TI Rapid fixation of disperse dyes on synthetic polymers
- L21 ANSWER 72 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN DUPLICATE 10  
TI Displacement of organic liquid films from solid surfaces by nonaqueous systems
- L21 ANSWER 73 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Exhaust dyeing of synthetic polymers with dyes dispersed in solution or emulsion in a saturated liquid fluorocarbon
- L21 ANSWER 74 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Perfluorinated ethers
- L21 ANSWER 75 OF 108 USPATFULL on STN  
TI Process for preparing perfluorinated ethers
- L21 ANSWER 76 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Colloidal-chemical properties of solutions of surfactants based on perfluoropropylene oxide oligomers. 1. Surface activity of ammonium salts of perfluoroolestermonocarboxylic acids at the aqueous solution-air interface
- L21 ANSWER 77 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Oligomeric fluorinated additives as surface modifiers for solid polymers
- L21 ANSWER 78 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN DUPLICATE 11  
TI Exhaust disperse dyeing of synthetic polymers using a saturated liquid fluorocarbon
- L21 ANSWER 79 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN DUPLICATE 12  
TI Bis-triazine compounds
- L21 ANSWER 80 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Exhaust disperse dyeing of synthetic fibers
- L21 ANSWER 81 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Fluoroalkylene ether difunctional compounds
- L21 ANSWER 82 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Dyeing synthetic fabrics with disperse dyes in fluorocarbon solvents
- L21 ANSWER 83 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Dispersion of dye in a fluorocarbon for exhaust dyeing
- L21 ANSWER 84 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN

10/630,698

- TI Rapid fixation of disperse dyes on synthetic polymers
- L21 ANSWER 85 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Fluorocarbon-dye dispersion for exhaust dispersion dyeing
- L21 ANSWER 86 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Exhaustion dyeing of films, fibers, and textiles of synthetic polymers with disperse dyes
- L21 ANSWER 87 OF 108 USPATFULL on STN  
TI Acrylic and methacrylic monomers, polymers and copolymers
- L21 ANSWER 88 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Perfluorinated linear polyethers having reactive terminal groups at both ends of the chain
- L21 ANSWER 89 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Perfluoroalkyletheramidoalkyl betaines and sulfobetaines
- L21 ANSWER 90 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI  $\alpha,\omega$ -Di-s-triazinyl perfluoroalkanes
- L21 ANSWER 91 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Polyfluoroalkoxy alkyl amidocarboxylic acids and salts
- L21 ANSWER 92 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Solid lubricant additives dispersed in perfluoroalkyl ethers with perfluoroalkyl ether acid dispersants
- L21 ANSWER 93 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Perfluoropolyethers by photooxidation of fluoroolefins
- L21 ANSWER 94 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Perfluoropoly(ether esters) as lubricants and hydraulic fluids
- L21 ANSWER 95 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Esters of hexafluoropropylene oxide polymer acids and polyalkylene glycols
- L21 ANSWER 96 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Liquid phase decarbonylation of fluorinated acyl fluorides
- L21 ANSWER 97 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Acrylate-type esters of perfluoropolyoxaalkaneamidoalkyl alcohols, and their polymers which are useful as oil and water repellents and as metal corrosion inhibitors
- L21 ANSWER 98 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Elastomers comprising acrylic and methacrylic derivatives of polyfluoropolyethers
- L21 ANSWER 99 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Polyfluoropolyoxaalkyl acrylates and N-(polyfluoropolyoxaalkyl)acrylamides
- L21 ANSWER 100 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Perfluoroalkyl ether amidoamine oxides
- L21 ANSWER 101 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Oil repellent polyfluoropolyoxo-alkyl phosphates
- L21 ANSWER 102 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Vaporization and decomposition kinetics of candidate re-entry blackout suppressants in low-pressure flames

- L21 ANSWER 103 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
 TI High temperature study of electrophilic gases for plasma quenching
- L21 ANSWER 104 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
 TI Corrosion-inhibited poly(hexafluoropropylene oxide) lubricants
- L21 ANSWER 105 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
 TI Perfluoro ketones
- L21 ANSWER 106 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
 TI Perfluorinated polyethers. Synthesis and characterization of a new class of inert fluids
- L21 ANSWER 107 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
 TI Fluorocarbon ethers from hexafluoropropylene oxide
- L21 ANSWER 108 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
 TI Dicarboxylic acids of fluorocarbon ethers and fluorides and their esters, amides, and salts

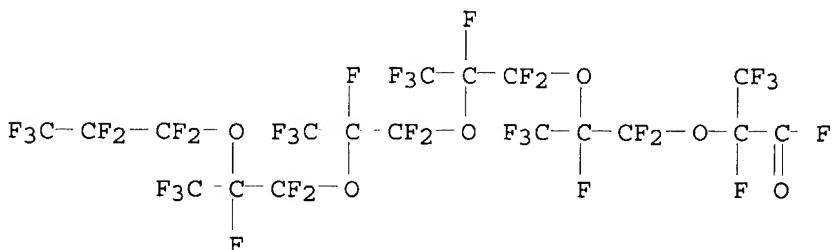
=> d 38,39,40,41,44,48,49,52,63,67,74,75,81,88,89,100,106 bib ab fhitstr

- L21 ANSWER 38 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
 AN 1989:614127 CAPLUS  
 DN 111:214127  
 TI Process for preparation of perfluorinated carboxylic acid fluorides  
 IN Kruse, Alfred; Siegemund, Guenter; Schwertfeger, Werner  
 PA Hoechst A.-G., Fed. Rep. Ger.  
 SO Ger. Offen., 5 pp.  
 CODEN: GWXXBX  
 DT Patent  
 LA German  
 FAN.CNT 1
- |      | PATENT NO.   | KIND | DATE     | APPLICATION NO. | DATE     |
|------|--|------|----------|-----------------|----------|
| PI   | DE 3737920   | A1   | 19890518 | DE 1987-3737920 | 19871107 |
|      | US 4874557   | A    | 19891017 | US 1988-266919  | 19881103 |
|      | EP 315908  | A1   | 19890517 | EP 1988-118391  | 19881104 |
|      | EP 315908  | B1   | 19920826 |                 |          |
|      | R: BE, CH, DE, FR, GB, IT, LI, NL  |      |          |                 |          |
|      | JP 01157933  | A2   | 19890621 | JP 1988-277536  | 19881104 |
|      | CN 1034199   | A    | 19890726 | CN 1988-107738  | 19881107 |
|      | CN 1022240   | B    | 19930929 |                 |          |
| PRAI | DE 1987-3737920  |      | 19871107 |                 |          |
| OS   | MARPAT 111:214127  |      |          |                 |          |
| AB   | F <sub>3</sub> CCF <sub>2</sub> [CF <sub>2</sub> OCF(CF <sub>3</sub> ) <sub>n</sub> COF (I; n = 2, 3), useful intermediates and monomers, are prepared by oligomerization of hexafluoropropylene oxide (II) at -20 to +100° in the presence of a catalyst system comprising: 1) alkali fluoride, preferably KF, 2-30%; 2) C <sub>5</sub> -8 alkanedinitrile, preferably adiponitrile, 50-95%; 3) MeO(CH <sub>2</sub> CH <sub>2</sub> O) <sub>m</sub> Me (III; m = 2-6, preferably 3) 2-50%. The process is advantageous in that higher temps. are used, product composition can be controlled by manipulation of the catalyst system composition, the product is readily separated, and the catalyst system can be reused. Thus, in a stainless steel autoclave a mixture of 30 g KF, 500 mL adiponitrile, and 100 mL III (m = 3) was stirred 30 min., continuously pressurized to 3.5 bar by addition of 5 kg II and stirred 2.5 h at 35-40°. After 3 h addnl. stirring the mixture readily separated into 2 phases. The lower product phase (4.90 kg) was drawn off and comprised the following I: n = 1, 21.5; n = 2, 61.1; n = 3, 16.3; and n = 4, 0.8%. |      |          |                 |          |
| IT   | 13252-15-8P  |      |          |                 |          |

RL: PREP (Preparation)

(manufacture of, by oligomerization of hexafluoropropylene oxide, catalysts for)

RN 13252-15-8 CAPLUS

CN 3,6,9,12,15-Pentaoxaoctadecanoyl fluoride, 2,4,4,5,7,7,8,10,10,11,13,13,14  
,16,16,17,17,18,18,18-eicosafuoro-2,5,8,11,14-pentakis(trifluoromethyl)-  
(7CI, 8CI, 9CI) (CA INDEX NAME)

L21 ANSWER 39 OF 108 USPATFULL on STN

AN 89:85709 USPATFULL

TI Process for the preparation of perfluorinated carbonyl fluorides

IN Kruse, Alfred, Kelkheim, Germany, Federal Republic of  
Siegemund, Gunter, Hofheim am Taunus, Germany, Federal Republic of  
Schwertfeger, Werner, Langgons, Germany, Federal Republic ofPA Hoechst Aktiengesellschaft, Frankfurt am Main, Germany, Federal Republic  
of (non-U.S. corporation)

PI US 4874557 19891017

AI US 1988-266919 19881103 (7)

PRAI DE 1987-3737920 19871107

DT Utility

FS Granted

EXNAM Primary Examiner: Killos, Paul J.

CLMN Number of Claims: 8

ECL Exemplary Claim: 1

DRWN No Drawings

LN.CNT 238

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The invention relates to a process for the preparation of perfluorinated carbonyl fluorides of the formula ##STR1## by oligomerization of hexafluoropropene oxide in the presence of a catalyst. The catalyst comprises a mixture of an alkali metal fluoride, a carboxylic acid dinitrile and a polyethylene glycol dimethyl ether.

IT 13252-15-8P

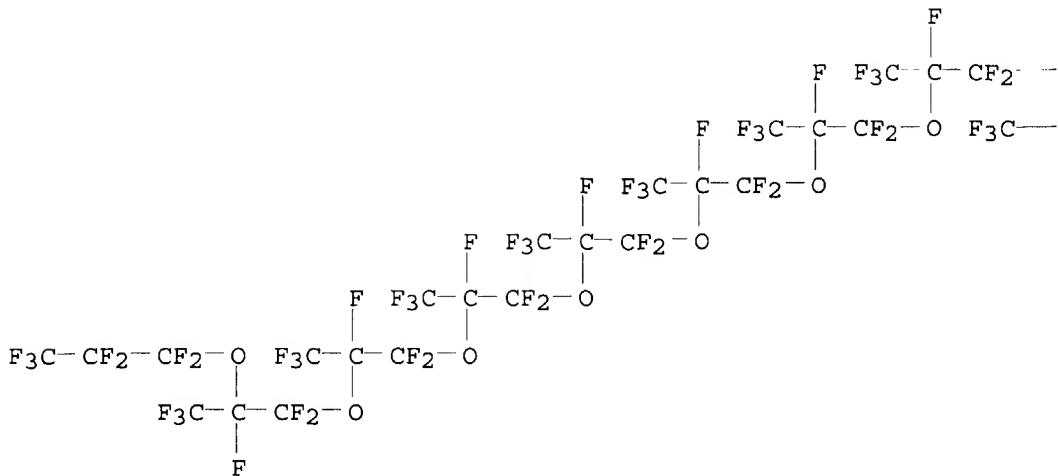
(manufacture of, by oligomerization of hexafluoropropylene oxide, catalysts for)

RN 13252-15-8 USPATFULL

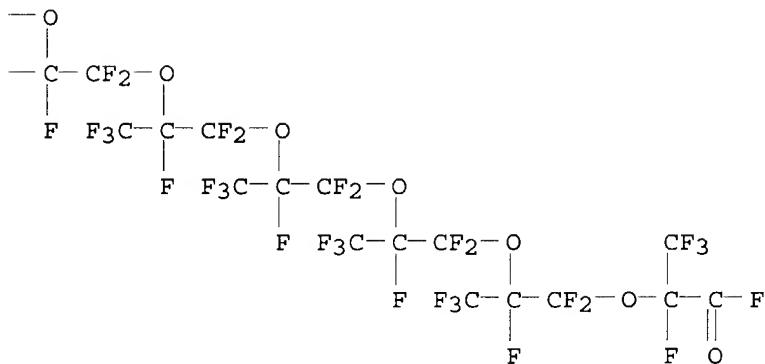
CN 3,6,9,12,15-Pentaoxaoctadecanoyl fluoride, 2,4,4,5,7,7,8,10,10,11,13,13,14  
,16,16,17,17,18,18,18-eicosafuoro-2,5,8,11,14-pentakis(trifluoromethyl)-  
(7CI, 8CI, 9CI) (CA INDEX NAME)



PAGE 1-A



PAGE 1-B



L21 ANSWER 41 OF 108 USPATFULL on STN

AN 88:5708 USPATFULL

TI Fluoropolyethers containing end groups endowed with anchoring capacity

IN Caporiccio, Gerardo, Milan, Italy

Strepparola, Ezio, Bergamo, Italy

Scarati, Mario A., Milan, Italy

PA Montedison S.p.A., Milan, Italy (non-U.S. corporation)

PI US 4721795 19880126

AI US 1984-687844 19841231 (6)

PRAI IT 1984-21481 19840619

DT Utility

FS Granted

EXNAM Primary Examiner: Chan, Nicky

LREP Stevens, Davis, Miller &amp; Mosher

CLMN Number of Claims: 3

ECL Exemplary Claim: 1

DRWN No Drawings

LN.CNT 442

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

10/630,698

AB Compounds suitable for being used as lubricants, having general formula:

(I) RO--(C.<sub>sub.3</sub> F.<sub>sub.6</sub> O).<sub>sub.m</sub>--(CFXO).<sub>sub.n</sub>--CFX--L, or

(II) R"CFXO--(C.<sub>sub.3</sub> F.<sub>sub.6</sub> O).<sub>sub.x</sub>(CFXO).<sub>sub.y</sub>--(C.<sub>sub.2</sub> F.<sub>sub.4</sub> O).<sub>sub.z</sub>--(CFX--L, where

R=--CF.<sub>sub.3</sub>, --C.<sub>sub.2</sub> F.<sub>sub.5</sub>, --C.<sub>sub.3</sub> F.<sub>sub.7</sub>

X=F, --CF.<sub>sub.3</sub>

R"=F, --CF.<sub>sub.3</sub>, --C.<sub>sub.2</sub> F.<sub>sub.5</sub>

m=an integer from 3 to 100

n=a finite integer, or=zero, wherefore m+n ranges from 3 to 100, provided that, if n is finite, m/n ranges from 5 to 20 and R is preferably=CF.<sub>sub.3</sub>, if n=zero, R is preferably --C.<sub>sub.2</sub> F.<sub>sub.5</sub> or --C.<sub>sub.3</sub> F.<sub>sub.7</sub>

x=a finite integer, or=zero

y, z=finite integers, such that x+y+z ranges from 5 to 200, while x+z/y ranges from 5 to 0.5, provided that when x=zero, z/y ranges from 1 to 0.5 and y+z ranges from 5 to 200 n while X is preferably F, and R"=L

L=group A--Y, where

A=--CH.<sub>sub.2</sub>O--, --CH.<sub>sub.2</sub>--O--CH.<sub>sub.2</sub>, --CF.<sub>sub.2</sub>, CF.<sub>sub.2</sub>O--,

Y=an organic radical covered by one of the following formulas: ##STR1## where R.<sub>sub.1</sub>, R.<sub>sub.2</sub>=alkyls C.<sub>sub.1</sub>-C.<sub>sub.3</sub>,

E=CHR.<sub>sub.3</sub> or --CH.<sub>sub.2</sub>--CHR.<sub>sub.3</sub>

B=H or a radical OR.<sub>sub.3</sub>--

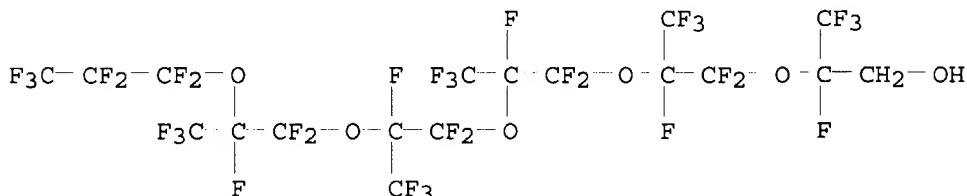
R.<sub>sub.3</sub>=H or an alkyl C.<sub>sub.1</sub>-C.<sub>sub.3</sub>.

IT 27617-34-1

(etherification of, by methylenedioxybenzyl chloride)

RN 27617-34-1 USPATFULL

CN 3,6,9,12,15-Pentaoxaoctadecan-1-ol, 2,4,4,5,7,7,8,10,10,11,13,13,14,16,16, 17,17,18,18,18-eicosfluoro-2,5,8,11,14-pentakis(trifluoromethyl)- (8CI, 9CI) (CA INDEX NAME)



L21 ANSWER 44 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN

AN 1986:592200 CAPLUS

DN 105:192200

TI Fluorine-containing polymers with oxygen permeability for medical use

IN Yamauchi, Koichi; Inoue, Yoshihisa; Yokoyama, Kazumasa

PA Green Cross Corp., Japan

10/630,698

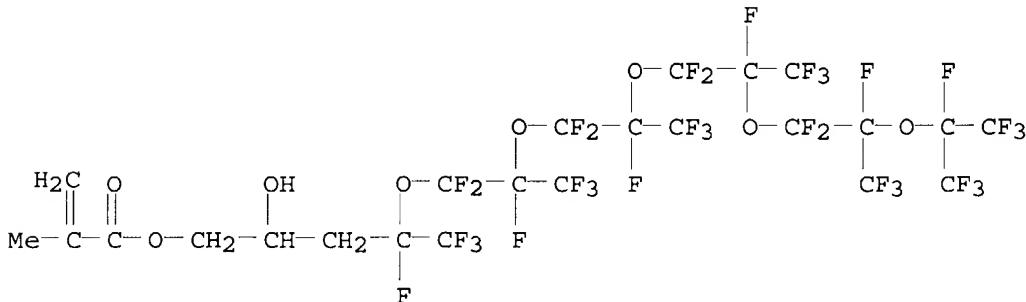
SO Jpn. Kokai Tokkyo Koho, 8 pp.  
CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 61111308	A2	19860529	JP 1984-233449	19841106
	JP 05061283	B4	19930906		
PRAI	JP 1984-233449		19841106		
AB	The title polymers, useful for hard contact lenses, were prepared from Y(CFXCF2O)mCFX'CH2CH(OH)CH2(OCH2CH2)nO2CCMe:CH2 (X, X' = F, lower perfluoroalkyl; Y = F, lower perfluoroalkoxy; m = 1-8; n = 0, 1) and have number-average mol. weight 700-20,000. Thus, (CF3)2CFO[CF(CF3)CF2O]4CF(CF3)Q				
(Q =	(glycidyl) was copolymerd. with trifluoroethyl methacrylate 0.50, vinylpyrrolidone 0.50, Me methacrylate 1.00, benzyl methacrylate 0.80, and allyl methacrylate 0.20 g in the presence of AIBN at 50° for 48 h, at 70° for 5 h, nd then at 90° for 3 h to obtain a button which was then dried at 110° for 2 days in vacuo to give Vicat hardness 11 and O permeation 21 + 10-11 cm3-cm/cm2-s-mm Hg.				
IT	104937-28-2P			RL: IMF (Industrial manufacture); PREP (Preparation) (manufacture and polymerization of)	
RN	104937-28-2 CAPLUS				
CN	2-Propenoic acid, 2-methyl-, 4,6,6,7,9,9,10,12,12,13,15,15,16,18,19,19,19-heptadecafluoro-2-hydroxy-4,7,10,13,16,18-hexakis(trifluoromethyl)-5,8,11,14,17-pentaoxanonadec-1-yl ester (9CI) (CA INDEX NAME)				



L21 ANSWER 48 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
AN 1986:186986 CAPLUS  
DN 104:186986  
TI Fluoropolyether compounds  
IN Caporiccio, Gerardo; Strepparola, Ezio; Scarati, Mario Alberto  
PA Montedison S.p.A., Italy  
SO Eur. Pat. Appl., 23 pp.  
CODEN: EPXXDW

DT Patent

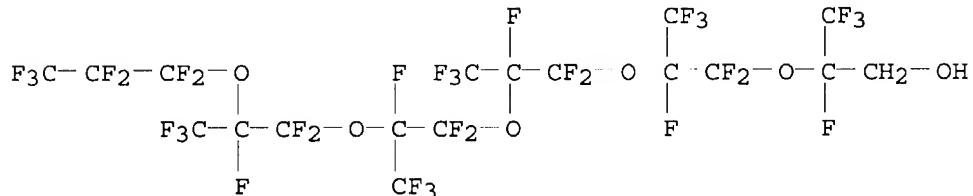
LA English

FAN.CNT 1

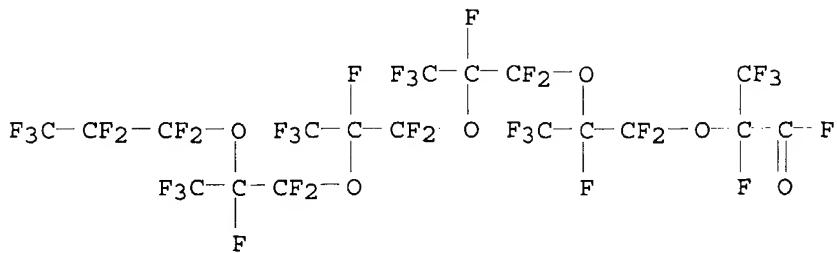
	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	EP 165650	A2	19851227	EP 1985-300785	19850206
	EP 165650	A3	19860219		
	EP 165650	B1	19890517		
	R: BE, DE, FR, GB, NL, SE				
	ES 539134	A1	19870501	ES 1984-539134	19841228

10/630,698

US 4721795 A 19880126 US 1984-687844 19841231  
 CA 1287637 A1 19910813 CA 1985-471406 19850103  
 JP 61004727 A2 19860110 JP 1985-3443 19850114  
 JP 06010257 B4 19940209  
 AU 8537757 A1 19860102 AU 1985-37757 19850117  
 AU 581640 B2 19890302  
 PRAI IT 1984-21481 19840619  
 AB Fluoro polyether compds. RO(C<sub>3</sub>F<sub>6</sub>O)<sub>m</sub>(CFXO)<sub>n</sub>CFXAY or R<sub>1</sub>CFXO(C<sub>3</sub>F<sub>6</sub>O)<sub>x</sub>(CFXO)<sub>y</sub>(C<sub>2</sub>F<sub>4</sub>O)<sub>z</sub>CFXAY (R = CF<sub>3</sub>, C<sub>2</sub>F<sub>5</sub>, C<sub>3</sub>F<sub>7</sub>; X = F, CF<sub>3</sub>; R<sub>1</sub> = F, CF<sub>3</sub>, C<sub>2</sub>F<sub>5</sub>; A = CH<sub>2</sub>O, CH<sub>2</sub>OCH<sub>2</sub>, CF<sub>2</sub>, CF<sub>2</sub>O; Y = dialkoxyphenyl, 1,2-methylenedioxyphenyl, etc.) are prepared for use as lubricants or protective coatings for audio or video tapes, floppy disks, etc. Thus, compound I was prepared from HOCH<sub>2</sub>CF<sub>2</sub>O(C<sub>2</sub>F<sub>4</sub>O)<sub>m</sub>(CF<sub>2</sub>O)<sub>p</sub>CF<sub>2</sub>CH<sub>2</sub>OH (m + p = 25, m/p = 0.6, mol. weight = 2300) 75, tert-BuOK 8, and 4-chloromethyl-1,2-methylenedioxybenzene 13 g. A 1% solution of I in Cl<sub>2</sub>CFCF<sub>2</sub>Cl was coated on a magnetic tape with CrO<sub>2</sub> pigment. The coating survived 8000 passages of a steel ball (diameter 0.32 mm) with 28 g load in an abrasion test, vs. 350 for a nonlubricated tape.  
 IT 27617-34-1  
 RL: RCT (Reactant); RACT (Reactant or reagent)  
 (etherification of, by methylenedioxybenzyl chloride)  
 RN 27617-34-1 CAPLUS  
 CN 3,6,9,12,15-Pentaoxaoctadecan-1-ol, 2,4,4,5,7,7,8,10,10,11,13,13,14,16,16,17,17,18,18,18-eicosfluoro-2,5,8,11,14-pentakis(trifluoromethyl)-(8CI, 9CI) (CA INDEX NAME)



L21 ANSWER 49 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
 AN 1984:474770 CAPLUS  
 DN 101:74770  
 TI Surface active substances containing an oligo(hexafluoropropene oxide) chain as a hydrophobic oleophobic moiety  
 AU Ishikawa, Nobuo; Sasabe, Mikio  
 CS Dep. Chem. Technol., Tokyo Inst. Technol., Tokyo, 152, Japan  
 SO Journal of Fluorine Chemistry (1984), 25(2), 241-53  
 CODEN: JFLCAR; ISSN: 0022-1139  
 DT Journal  
 LA English  
 AB Oil-soluble surfactants  $\text{CF}_3\text{CF}_2\text{CF}_2\text{O}[\text{CF}(\text{CF}_3)\text{CF}_2\text{O}]n-2\text{CF}(\text{CF}_3)\text{COR}$  (I) ( $\text{R} = \text{Ph}$  or  $p\text{-tolyl}$ ,  $n = 2-6$ ) were prepared by acylating arenes with hexafluoropropylene oxide oligomers. These surfactants (0.2-0.5%) decreased the surface tensions of toluene and  $m\text{-xylene}$  to 12-14 dynes/cm. Water-soluble surfactants I [ $\text{R} = m\text{-(NaO}_3\text{S)}\text{C}_6\text{H}_4$  or  $4\text{-Me-3-(NaO}_3\text{S)}\text{C}_6\text{H}_3$ ,  $n = 2-6$ ] were also prepared. Some of the surfactants (i.e.,  $n = 4-6$ ) decreased the surface tension of water to 16 dynes/cm at a concentration of 10-4-10-5M.  
 IT 13252-15-8  
 RL: RCT (Reactant); RACT (Reactant or reagent)  
 (acylation by, of arenes)  
 RN 13252-15-8 CAPLUS  
 CN 3,6,9,12,15-Pentaoxaoctadecanoyl fluoride, 2,4,4,5,7,7,8,10,10,11,13,13,14,16,16,17,17,18,18,18-eicosfluoro-2,5,8,11,14-pentakis(trifluoromethyl)-(7CI, 8CI, 9CI) (CA INDEX NAME)



L21 ANSWER 52 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
AN 1983:145452 CAPLUS

DN 98:145452

TI Synthesis of fluorinated surfactants containing hexafluoropropene oxide as a hydrophobic group and properties of the solutions

AU Ogino, Keizo; Murakami, Hiroki; Ishikawa, Nobuo; Sasabe, Mikio

CS Fac. Sci. Technol., Sci. Univ. Tokyo, Noda, Japan

SO Yukagaku (1983), 32(2), 96-101

CODEN: YKGKAM; ISSN: 0513-398X

DT Journal

LA Japanese

AB The surfactants  $C_3F_7O[CF(CF_3)CF_2O]n-2CF(CF_3)CO_2Na$  (I) ( $n = 2-6$ ) were prepared. The critical micelle concentration decreases with increasing  $n$ . A secondary

critical micelle concentration is observed for I ( $n = 4-6$ ). The Krafft points of I are

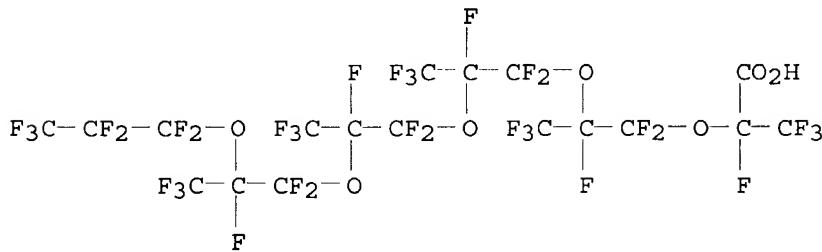
$<0^\circ$ . I ( $n = 4$ ) [67963-78-4] has the best foaming properties. I are stable in acidic and alkaline solns.

IT 85248-41-5P

RL: SPN (Synthetic preparation); PREP (Preparation)  
(preparation and surfactant properties of)

RN 85248-41-5 CAPLUS

CN 3,6,9,12,15-Pentaoxaocadecanoic acid, 2,4,4,5,7,7,8,10,10,11,13,13,14,16,  
16,17,17,18,18,18-eicosafuoro-2,5,8,11,14-pentakis(trifluoromethyl)-,  
sodium salt (9CI) (CA INDEX NAME)



● Na

L21 ANSWER 63 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN

AN 1978:508039 CAPLUS

DN 89:108039

TI Synthesis of perfluoro(polyether) difunctional compounds

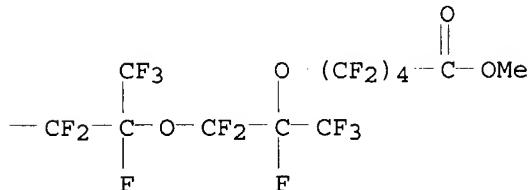
AU Soloski, E. J.; Tamborski, C.; Psarras, T.

10/630,698

CS Air Force Mater. Lab., Wright-Patterson AFB, OH, USA  
SO Journal of Fluorine Chemistry (1978), 11(6), 601-12  
CODEN: JFLCAR; ISSN: 0022-1139  
DT Journal  
LA English  
AB  $\omega$ -Iodoperfluoro(polyether) esters IRFOQfCO<sub>2</sub>R (I; Rf = perfluoroalkylene, Qf = perfluoroalkylene moiety containing O atoms in chain, R = Me or Et) were prepared by 2 procedures. I reacted via Zn coupling reactions to give  $\alpha$ , $\omega$ -perfluoro(polyether) diesters. The diesters serve as convenient starting materials for the preparation of a variety of other difunctional compds. of high mol. weight and exhibiting a variation of O-C ratio.  
IT 61210-96-6P  
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)  
(preparation and amidation of)  
RN 61210-96-6 CAPLUS  
CN 6,9,12,15,20,23,26,29-Octaoxatetratriacontanedioic acid,  
2,2,3,3,4,4,5,5,7,8,8,10,11,11,13,14,14,16,16,17,17,18,18,19,19,21,21,22,2  
4,24,25,27,27,28,30,30,31,31,32,32,33,33-dotetracontafluoro-  
7,10,13,22,25,28-hexakis(trifluoromethyl)-, dimethyl ester (9CI) (CA  
INDEX NAME)

\*\*\* STRUCTURE DIAGRAM IS NOT AVAILABLE \*\*\*

PAGE 1-B



L21 ANSWER 67 OF 108 USPATFULL on STN  
AN 77:11582 USPATFULL  
TI Fluoroalkyleneether difunctional compounds  
IN Tamborski, Christ, Dayton, OH, United States  
PA The United States of America as represented by the Secretary of the Air Force, Washington, DC, United States (U.S. government)  
PI US 4011255 19770308  
AI US 1975-610520 19750904 (5)  
DT Utility  
FS Granted  
EXNAM Primary Examiner: Brust, Joseph Paul  
LREP Rusz, Joseph E., Kuhn, Cedric H.  
CLMN Number of Claims: 3  
ECL Exemplary Claim: 1  
DRWN No Drawings  
LN.CNT 304  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.  
AB Omega-carbomethoxyperfluoroalkylene ether iodides are reacted with metallic zinc to yield alpha-omega perfluoroalkyleneether diesters. The diesters are reacted with ammonia to form diamides, the diamides are reacted with phosphorus pentoxide to form dinitriles, and the dinitriles are esterified with methanol to form diimide esters. The diimide esters are particularly useful as monomers in synthesizing

10/630,698

perfluoroalkylene ether bibenzoxazole polymers possessing  
thermooxidative stability and outstanding low temperature viscoelastic  
properties.

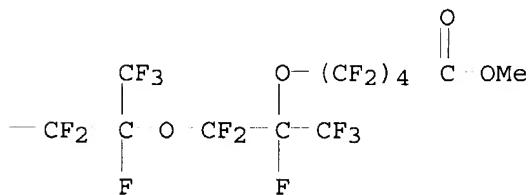
IT 61210-96-6P  
(preparation and amidation of)

RN 61210-96-6 USPATFULL

CN 6,9,12,15,20,23,26,29-Octaoxatetratriacontanedioic acid,  
2,2,3,3,4,4,5,5,7,8,8,10,11,11,13,14,14,16,16,17,17,18,18,19,19,21,21,22  
,24,24,25,27,27,28,30,30,31,31,32,32,33,33-dotetracontafluoro-  
7,10,13,22,25,28-hexakis(trifluoromethyl)-, dimethyl ester (9CI) (CA  
INDEX NAME)

\*\*\* STRUCTURE DIAGRAM IS NOT AVAILABLE \*\*\*

PAGE 1-B



L21 ANSWER 74 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN

AN 1976:463699 CAPLUS

DN 85:63699

TI Perfluorinated ethers

IN Von Halasz, Sigmar P.; Kluge, Friedhelm

PA Hoechst A.-G., Fed. Rep. Ger.

SO Ger. Offen., 24 pp.

CODEN: GWXXBX

DT Patent

LA German

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	DE 2451493	A1	19760506	DE 1974-2451493	19741030
	DE 2451493	C2	19820624		
	NL 7512495	A	19760504	NL 1975-12495	19751024
	US 3985810	A	19761012	US 1975-626349	19751028
	GB 1484823	A	19770908	GB 1975-44343	19751028
	CA 1060482	A1	19790814	CA 1975-238542	19751029
	FR 2289477	A1	19760528	FR 1975-33188	19751030
	FR 2289477	B1	19790105		

PRAI DE 1974-2451493 19741030

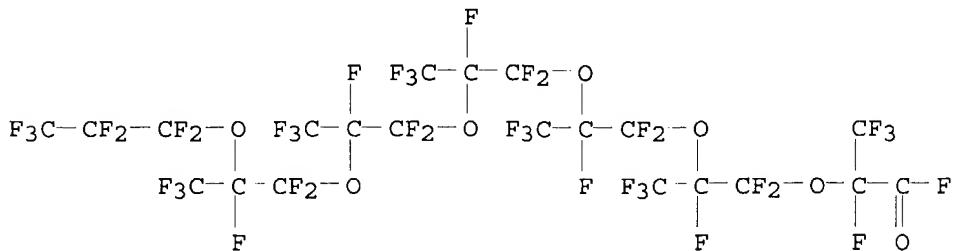
AB The polyethers  $\text{Rf}[(\text{OCF}(\text{R})\text{CF}_2)_x\text{OCF}_2\text{R}]_n$  (I) ( $\text{R} = \text{F}, \text{CF}_3$ ;  $\text{Rf} = \text{perfluoroalkyl}$  or  $\text{perfluoroalkylene}$ ;  $n = 1-2$ ;  $x = 0-50$ ), useful as hydraulic fluids, heat transfer media, lubricants, etc., are prepared by reaction of F with  $\text{Rf}[(\text{OCF}(\text{R})\text{CF}_2)_x\text{OCF}(\text{R})\text{COF}]_n$  (II) in the presence of metal catalysts at  $50-350^\circ$ . Thus, adding 439.5 g II ( $\text{R} = \text{CF}_3$ ,  $\text{Rf} = \text{CF}(\text{CF}_3)\text{CF}(\text{CF}_3)$ ,  $n = 2$ ,  $x = 6.5-9.5$ ) [59859-32-4] over 19.5 hr to a Cu tube packed with silvered Cu filings with countercurrent addition of 0.8 l./hr 3:1 F-He at  $200-5^\circ$  gives 405 g I ( $\text{R} = \text{CF}_3$ ,  $\text{Rf} = \text{CF}(\text{CF}_3)\text{CF}(\text{CF}_3)$ ,  $n = 2$ ,  $x = 13.5-19.5$ ) [59859-33-5], b0.4-0.5 185-280°.

IT 13140-24-4

RL: RCT (Reactant); RACT (Reactant or reagent)  
(fluorination of, to perfluoroalkyl ethers)

10/630,698

RN 13140-24-4 CAPPLUS  
CN 3,6,9,12,15,18-Hexaoxaheneicosanoyl fluoride,  
2,4,4,5,7,7,8,10,10,11,13,13,14,16,16,17,19,19,20,20,21,21,21-  
tricosfluoro-2,5,8,11,14,17-hexakis(trifluoromethyl)- (7CI, 8CI, 9CI)  
(CA INDEX NAME)



L21 ANSWER 75 OF 108 USPATFULL on STN

AN 76:55823 USPATFULL

TI Process for preparing perfluorinated ethers

IN von Halasz, Sigmar-Peter, Kelkheim, Taunus, Germany, Federal Republic of Kluge, Friedhelm, Frankfurt am Main, Germany, Federal Republic of

PA Hoechst Aktiengesellschaft, Frankfurt am Main, Germany, Federal Republic of (non-U.S. corporation)

PI US 3985810 19761012

AI US 1975-626349 19751028 (5)

PRAI DE 1974-2451493 19741030

DT Utility

FS Granted

EXNAM Primary Examiner: Mars, Howard T.

LREP Curtis, Morris & Safford

CLMN Number of Claims: 10

ECL Exemplary Claim: 1

DRWN 1 Drawing Figure(s); 1 Drawing Page(s)

LN.CNT 600

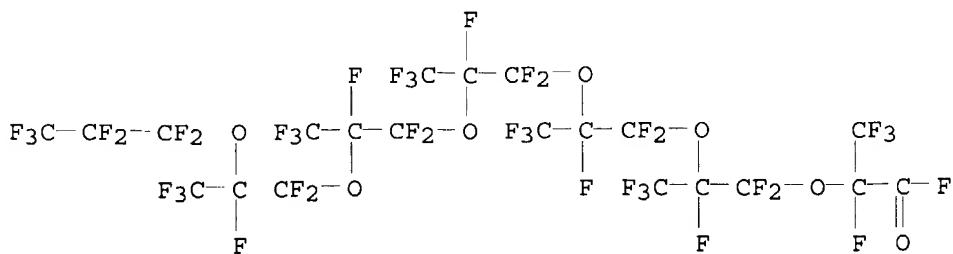
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Perfluorinated ethers containing carboxylic acid fluoride groups and optionally units derived from hexafluoropropene epoxide or tetrafluoroethylene epoxide are reacted with fluorine at temperatures of from 50° to 350°C in the presence of metallic catalysts. During the reaction carbonyl difluoride is split off and an ether is obtained in high yield which is free of carboxylic acid fluoride groups. Metallic silver is well suited as catalyst.

IT 13140-24-4  
(fluorination of, to perfluoroalkyl ethers)

RN 13140-24-4 USPATFULL

CN 3,6,9,12,15,18-Hexaoxaheneicosanoyl fluoride,  
2,4,4,5,7,7,8,10,10,11,13,13,14,16,16,17,19,19,20,20,21,21,21-  
tricosfluoro-2,5,8,11,14,17-hexakis(trifluoromethyl)- (7CI, 8CI, 9CI)  
(CA INDEX NAME)



L21 ANSWER 81 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN

AN 1977:4951 CAPLUS

DN 86:4951

TI Fluoroalkylene ether difunctional compounds

IN Tamborski, Christ

PA United States Dept. of the Air Force, USA

SO U. S. Pat. Appl., 14 pp. Avail. NTIS.

CODEN: XAXXAV

DT Patent

LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 610520	A0	19750904	US 1975-610520	19750904
PRAI	US 1975-610520		19750904		

AB MeOC(:NH)(CF<sub>2</sub>)<sub>4</sub>O(CF<sub>2</sub>)<sub>4</sub>O(:NH)OMe, a monomer for preparation of elastomeric thermally stable polymers, was prepared by treating ICF<sub>2</sub>CF<sub>2</sub>O(CF<sub>2</sub>)<sub>4</sub>CO<sub>2</sub>Et (I) with Zn to give EtO<sub>2</sub>C(CF<sub>2</sub>)<sub>4</sub>O(CF<sub>2</sub>)<sub>4</sub>CO<sub>2</sub>Et, treatment of the diester with NH<sub>3</sub> to give the diamide, dehydration of the diamide with P<sub>2</sub>O<sub>5</sub> to give the dinitrile, and treatment of the latter with Na-MeOH to give the diimide. MeO<sub>2</sub>C(CF<sub>2</sub>)<sub>4</sub>[CF(CF<sub>3</sub>)CF<sub>2</sub>O]<sub>n</sub>CF<sub>2</sub>CF<sub>2</sub>I (n = 2, 3) and MeO<sub>2</sub>CCF(CF<sub>3</sub>)OCF<sub>2</sub>CF<sub>2</sub>OCF(CF<sub>3</sub>)CF<sub>2</sub>OCF<sub>2</sub>CF<sub>2</sub>I were also used in place of I.

IT 61210-96-6P

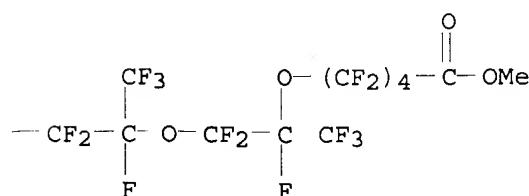
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)  
(preparation and amidation of)

RN 61210-96-6 CAPLUS

CN 6,9,12,15,20,23,26,29-Octaoxatetracontanedioic acid,  
2,2,3,3,4,4,5,5,7,8,8,10,11,11,13,14,14,16,16,17,17,18,18,19,19,21,21,22,2  
4,24,25,27,27,28,30,30,31,31,32,32,33,33-dotetracontafluoro-  
7,10,13,22,25,28-hexakis(trifluoromethyl)-, dimethyl ester (9CI) (CA  
INDEX NAME)

\*\*\* STRUCTURE DIAGRAM IS NOT AVAILABLE \*\*\*

PAGE 1-B



10/630,698

L21 ANSWER 88 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
AN 1975:594115 CAPLUS  
DN 83:194115

TI Perfluorinated linear polyethers having reactive terminal groups at both ends of the chain

IN Sianesi, Dario; Caporiccio, Gerardo; Mensi, Domenico

PA Montedison S.p.A., Italy

SO U.S., 14 pp.

CODEN: USXXAM

DT Patent

LA English

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI US 3847978	A	19741112	US 1969-834486	19690618
PRAI US 1968-787309		19681226		

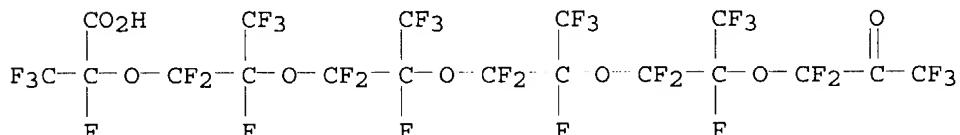
AB Perfluorinated linear polyethers containing peroxidic linkages were chain-cleaved by reducing agents to give bifunctional perfluorinated linear oligopolyethers with chemical-reactive terminal groups. Thus, hexafluoropropene [116-15-4] was treated with oxygen under the influence of uv light to give a peroxidized poly(perfluoropropylene oxide) [25038-02-2] which was reduced by H over a Pd catalyst to give a series of carboxy- and trifluoroacetyl-terminated oligopolyethers. One of these, CF<sub>3</sub>COCF<sub>2</sub>O(C<sub>3</sub>F<sub>6</sub>O)<sub>2</sub>CF(CF<sub>3</sub>)CO<sub>2</sub>H [42775-40-6], boiling point 210-2°, formed a polymer with hexamethylenediamine [55809-69-3].

IT 42775-42-8P

RL: IMF (Industrial manufacture); PREP (Preparation)  
(manufacture of, by reduction of perfluorinated polyether peroxy derivs.)

RN 42775-42-8 CAPLUS

CN 3,6,9,12,15-Pentaoxaoctadecanoic acid, 2,4,4,5,7,7,8,10,10,11,13,13,14,16,  
16,18,18,18-octadecafluoro-17-oxo-2,5,8,11,14-pentakis(trifluoromethyl)-  
(9CI) (CA INDEX NAME)



L21 ANSWER 89 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN

AN 1975:88056 CAPLUS

DN 82:88056

TI Perfluoroalkyletheramidoalkyl betaines and sulfobetaines

IN Barlett, Phillip Lee

PA du Pont de Nemours, E. I., and Co.

SO U.S., 4 pp.

CODEN: USXXAM

DT Patent

LA English

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI US 3839425	A	19741001	US 1970-72803	19700916
PRAI US 1970-72803		19700916		

AB Thirteen surfactants [C<sub>3</sub>F<sub>7</sub>O[CF(CF<sub>3</sub>)CF<sub>2</sub>O]<sub>n</sub>CF(CF<sub>3</sub>)CONR(CH<sub>2</sub>)<sub>m</sub>N<sup>+</sup>(R<sub>1</sub>)<sub>2</sub>R<sub>2</sub> with R = H, Me, or Et, R<sub>1</sub> = Me, Et, or Pr, R<sub>2</sub> = (CH<sub>2</sub>)<sub>1-2</sub>CO<sub>2</sub>- or (CH<sub>2</sub>)<sub>2-3</sub>SO<sub>3</sub>-, m = 2-3, and n = 0-4] were prepared and were especially useful as foam stabilizers for

10/630,698

fire-extinguishing foams on burning hydrocarbon surfaces. Thus, 50 g C<sub>3</sub>F<sub>7</sub>OCF(CF<sub>3</sub>)CF<sub>2</sub>OCF(CF<sub>3</sub>)CONH(CH<sub>2</sub>)<sub>3</sub>NMe<sub>2</sub> [31339-59-0], 10.1 g ClCH<sub>2</sub>CO<sub>2</sub>Na [3926-62-3], and 20 ml iso-PrOH were refluxed for 16 hr to prepare 53.7 g C<sub>3</sub>F<sub>7</sub>OCF(CF<sub>3</sub>)CF<sub>2</sub>OCF(CF<sub>3</sub>)CONH(CH<sub>2</sub>)<sub>3</sub>N+Me<sub>2</sub>CH<sub>2</sub>CO<sub>2</sub>- [54190-98-6] which gave surface tension 18.7 dynes/cm as a 0.001% aqueous solution and, as a 1% solution,

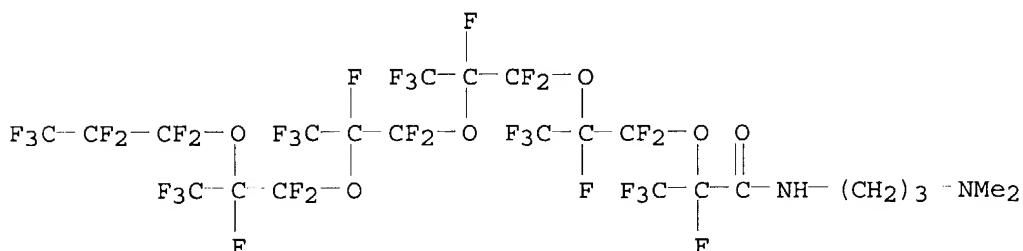
caused water to spread rapidly over the surface of cyclohexane.

IT 54190-86-2

RL: RCT (Reactant); RACT (Reactant or reagent)  
(reaction of, with carboxyalkyl and sulfoalkyl halides)

RN 54190-86-2 CAPLUS

CN 3,6,9,12,15-Pentaoxaoctadecanamide, N-[3-(dimethylamino)propyl]-  
2,4,4,5,7,7,8,10,10,11,13,13,14,16,16,17,17,18,18,18-eicosfluoro-  
2,5,8,11,14-pentakis(trifluoromethyl)- (9CI) (CA INDEX NAME)



L21 ANSWER 100 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN

AN 1972:16000 CAPLUS

DN 76:16000

TI Perfluoroalkyl ether amidoamine oxides

IN Bartlett, Philip L.

PA du Pont de Nemours, E. I., and Co.

SO U.S., 6 pp.

CODEN: USXXAM

DT Patent

LA English

FAN.CNT 3

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 3547995	A	19701215	US 1968-705932	19680216
	NL 6803275	A	19680909	NL 1968-3275	19680307
	FR 1568163	A	19690523	FR 1968-1568163	19680307
	GB 1202830	A	19700819	GB 1968-1202830	19680307
	DE 1793761	A1	19730823	DE 1967-1793761	19680307

PRAI	US 1967-621128	19670307
	US 1967-621148	19670307
	US 1967-621157	19670307
	US 1968-705923	19680216
	US 1968-705932	19680216
	US 1968-705947	19680216

AB A group of perfluoroalkyl ether amidoamine oxides are useful as surface active agents and are noncorrosive to steel. Hexafluoropropylene oxide is trimerized to perfluoroalkyl ether acid fluoride which is esterified with methanol and reacted with 3-(dimethylamino)propylamine to give a corresponding perfluoro alkyl ether amide which was oxidized to [3-[2-[2-(heptafluoropropoxy)hexafluoropropoxy]tetrafluoropropionamido]propyl]dimethylamine oxide [29209-86-7].

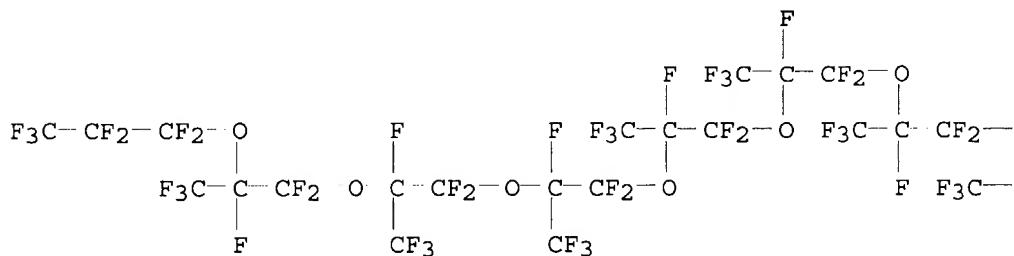
IT 34839-72-0

RL: TEM (Technical or engineered material use); USES (Uses)  
(surfactants)

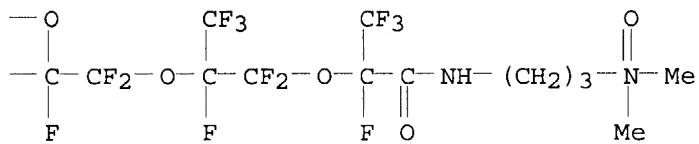
10/630,698

RN 34839-72-0 CAPLUS  
CN 3,6,9,12,15,18,21,24-Octaoxatriacontanamide, N-[3-(dimethyloxidoamino)propyl]-2,4,4,5,7,7,8,10,10,11,13,13,14,16,16,17,19,19,20,22,22,23,25,25,26,28,28,29,29,30,30,30-dotriacontafluoro-2,5,8,11,14,17,20,23,26-nonakis(trifluoromethyl)- (9CI) (CA INDEX NAME)

PAGE 1-A



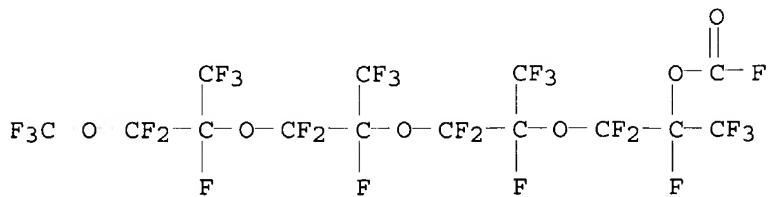
PAGE 1-B



L21 ANSWER 106 OF 108 CAPLUS COPYRIGHT 2004 ACS on STN  
AN 1968:77639 CAPLUS  
DN 68:77639  
TI Perfluorinated polyethers. Synthesis and characterization of a new class of inert fluids  
AU Sianesi, Dario  
CS "Montecatini Edison", Ist. "G. Donegani", Milan-Linate, Italy  
SO Chimica e l'Industria (Milan, Italy) (1968), 50(2), 206-14  
CODEN: CINMAB; ISSN: 0009-4315  
DT Journal  
LA Italian  
AB The photochem. reaction between hexafluoropropylene and O was studied. Compds. of the general formulas CF<sub>3</sub>(OR)<sub>n</sub>O[CF<sub>2</sub>CF(CF<sub>3</sub>)]<sub>m</sub>COF (I), CF<sub>3</sub>(OR)<sub>n</sub>O[CF<sub>2</sub>CF(CF<sub>3</sub>)]<sub>m</sub>CF<sub>2</sub>COCF<sub>3</sub> (II), and CF<sub>3</sub>(OR)<sub>n</sub>O[CF<sub>2</sub>CF(CF<sub>3</sub>)]<sub>m</sub>CF<sub>2</sub>H (III) are obtained. The following I (n, R, m, and b.p. given): 0, -, 0, -, 0, 51°; 0, -, 2, 114°; 0, -, 3, 156-7°; 0, -, 4, 195-7°; 1, CF<sub>2</sub>, 1, 83-6°; 1, CF<sub>2</sub>, 2, 130-3°; 1, CF(CF<sub>2</sub>), 1, 96-8°; 1, CF(CF<sub>3</sub>), 2, 143-5°; the following II (n, R, m, and b.p. given): 0, -, 0, 15°; 0, -, 1, 87°; 0, -, 2, 137°; 0, -, 3, 180-1°; 0, -, 4, 215-16°; 0, -, 5, 244-5°; 1, CF<sub>2</sub>, 0, 48-9°; 1, CF<sub>2</sub>, 1, 106-7°; 1, CF<sub>2</sub>, 2, 157-60°; 1, CF<sub>2</sub>, 3, 197-200°; 1, CF(CF<sub>3</sub>), 0, 68°; 1, CF(CF<sub>3</sub>), 1, 121-2°; 1, CF(CF<sub>3</sub>), 2, 168-70°; 1, CF(CF<sub>3</sub>), 3, 205-7°; and the following III (n, R, m, and b.p. given): 1, -, 0, -36°; 0, -, 1, 55°; 0, -, 2, 113°; 0, -, 3, 161-2°; 0, -, 4, 200-1°; 0, -, 5, 231-2°; 1, CF<sub>2</sub>, 1, 88-90°; 1, CF<sub>2</sub>, 2, 133-4°; 1, CF<sub>2</sub>, 3, 175-8°; 1, CF(CF<sub>3</sub>), 1, 100-1°; 1, CF(CF<sub>3</sub>), 2, 147-8°; 1, CF(CF<sub>3</sub>), 3, 189-92°, are prepared

10/630,698

IT 18934-94-6P  
RL: SPN (Synthetic preparation); PREP (Preparation)  
(preparation of)  
RN 18934-94-6 CAPLUS  
CN Formic acid, fluoro-, trifluoro-2-[trifluoro-1-(trifluoromethyl)-2-[trifluoro-1-(trifluoromethyl)-2-[trifluoro-2-(trifluoromethoxy)-1-(trifluoromethyl)ethoxy]ethoxy]-1-(trifluoromethyl)ethyl ester  
(8CI) (CA INDEX NAME)



	SINCE FILE ENTRY	TOTAL SESSION
COST IN U.S. DOLLARS		
FULL ESTIMATED COST	152.55	469.93
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LAST RELOADED: Nov 5, 2004 (20041105/UP).

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COST IN U.S. DOLLARS		
FULL ESTIMATED COST	0.18	470.11
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CA SUBSCRIBER PRICE	0.00	-9.10

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STRUCTURE FILE UPDATES: 8 NOV 2004 HIGHEST RN 777024-10-9  
DICTIONARY FILE UPDATES: 8 NOV 2004 HIGHEST RN 777024-10-9

TSCA INFORMATION NOW CURRENT THROUGH MAY 21, 2004

Please note that search-term pricing does apply when  
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10/630,698

Crossover limits have been increased. See HELP CROSSOVER for details.

Experimental and calculated property data are now available. For more information enter HELP PROP at an arrow prompt in the file or refer to the file summary sheet on the web at:  
<http://www.cas.org/ONLINE/DBSS/registryss.html>

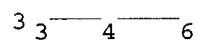
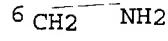
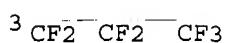
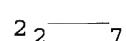
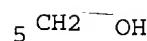
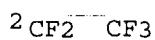
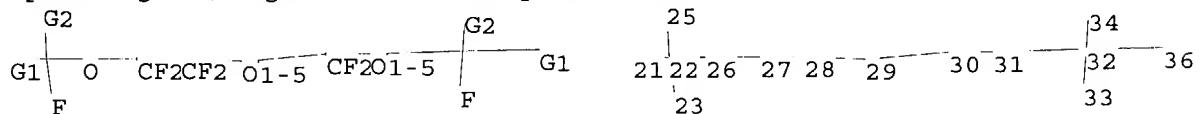
=> ....Testing the current file.... screen

ENTER SCREEN EXPRESSION OR (END) :end

=> screen 1994 OR 2016 OR 2021 OR 2026 OR 1838

L22 SCREEN CREATED

=>  
Uploading C:\Program Files\Stnexp\Queries\10630698b-1.str



chain nodes :  
1 2 3 4 5 6 7 8 9 10 11 12 21 22 23 25 26 27 28 29 30 31 32  
33 34 36  
chain bonds :  
2-7 3-4 4-6 8-10 9-11 21-22 22-23 22-25 22-26 26-27 27-28 28-29 29-30  
30-31 31-32 32-36 32-33 32-34  
exact/norm bonds :

10/630,698

21-22 22-25 22-26 31-32 32-36 32-34  
exact bonds :  
2-7 3-4 4-6 8-10 9-11 22-23 26-27 27-28 28-29 29-30 30-31 32-33

G1: [\*1], [\*2], [\*3], [\*4], [\*5], [\*6], [\*7]

G2: [\*1], [\*2]

Match level :

1:CLASS 2:CLASS 3:CLASS 4:CLASS 5:CLASS 6:CLASS 7:CLASS 8:CLASS 9:CLASS  
10:CLASS 11:CLASS 12:CLASS 21:CLASS 22:CLASS 23:CLASS 25:CLASS 26:CLASS  
27:CLASS 28:CLASS 29:CLASS 30:CLASS 31:CLASS 32:CLASS 33:CLASS 34:CLASS  
36:CLASS

L23 STRUCTURE UPLOADED

=> que L23 NOT L22

L24 QUE L23 NOT L22

=> s l24  
SAMPLE SEARCH INITIATED 07:31:27 FILE 'REGISTRY'  
SAMPLE SCREEN SEARCH COMPLETED - 265 TO ITERATE

100.0% PROCESSED 265 ITERATIONS ( 4 INCOMPLETE) 6 ANSWERS  
SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE \*\*COMPLETE\*\*  
BATCH \*\*COMPLETE\*\*  
PROJECTED ITERATIONS: 4324 TO 6276  
PROJECTED ANSWERS: 6 TO 266

L25 6 SEA SSS SAM L23 NOT L22

=> s l24 ful  
FULL SEARCH INITIATED 07:31:35 FILE 'REGISTRY'  
FULL SCREEN SEARCH COMPLETED - 4703 TO ITERATE

100.0% PROCESSED 4703 ITERATIONS ( 70 INCOMPLETE) 81 ANSWERS  
SEARCH TIME: 00.00.03

L26 81 SEA SSS FUL L23 NOT L22

=> file caplus casreact uspatful  
COST IN U.S. DOLLARS SINCE FILE TOTAL  
ENTRY SESSION  
FULL ESTIMATED COST 156.26 626.37  
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) SINCE FILE TOTAL  
ENTRY SESSION  
CA SUBSCRIBER PRICE 0.00 -9.10

FILE 'CAPLUS' ENTERED AT 07:32:23 ON 10 NOV 2004  
USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.  
PLEASE SEE "HELP USAGETERMS" FOR DETAILS.  
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10/630,698

USE IS SUBJECT TO THE TERMS OF YOUR CUSTOMER AGREEMENT  
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FILE 'USPATFULL' ENTERED AT 07:32:23 ON 10 NOV 2004  
CA INDEXING COPYRIGHT (C) 2004 AMERICAN CHEMICAL SOCIETY (ACS)

=> s 126  
L27 76 L26

=> dup rem 127  
PROCESSING COMPLETED FOR L27  
L28 69 DUP REM L27 (7 DUPLICATES REMOVED)

=> d 1-69 ti

L28 ANSWER 1 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Manufacture of magnetic recording media

L28 ANSWER 2 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Perfluoropoly-ether/peroxide compounds: spectroscopic studies and quantum chemical calculations

L28 ANSWER 3 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Fluorine-containing compounds, lubricants and magnetic recording media therewith, and manufacture thereof

L28 ANSWER 4 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Manufacture of magnetic recording media

L28 ANSWER 5 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Fluorine-containing tertiary amine tricarboxylate ester, lubricant, magnetic recording medium using the lubricant, and manufacture of the recording medium

L28 ANSWER 6 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Magnetic recording media having good traveling durability and high electromagnetic conversion and their manufacture

L28 ANSWER 7 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Perfluoroalkyl polyether oligomers containing phosphazene groups useful as lubricants for recording media such as hard disks

L28 ANSWER 8 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Magnetic recording medium and its fabrication

L28 ANSWER 9 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Thin magnetic tapes with good durability having stainless reinforcing layers on their back side and their manufacture

L28 ANSWER 10 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Magnetic tapes with fluorine-containing lubricant layers and their manufacture

L28 ANSWER 11 OF 69 USPATFULL on STN  
TI Rolling bearing

L28 ANSWER 12 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
TI (Fluoroorgano)silicon compounds as hydro- and oleophobic agents for protection of building materials from adverse effects of environment

L28 ANSWER 13 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Lubricating grease for sliding bearings

10/630,698

- L28 ANSWER 14 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Monoesters of fluorinated alkylidicarboxylic acids, lubricant compositions, magnetic recording media, and their manufacture
- L28 ANSWER 15 OF 69 USPATFULL on STN  
TI Magnetic recording medium having a perfluoropolyether lubricant bonded to the surface of a carbon protective film
- L28 ANSWER 16 OF 69 USPATFULL on STN  
TI Liquid-phase fluorination
- L28 ANSWER 17 OF 69 USPATFULL on STN  
TI Liquid-phase fluorination
- L28 ANSWER 18 OF 69 USPATFULL on STN  
TI Amides and esters of perfluoropolyoxaalkylene-sulfo- or perfluoropolyoxaalkylene-carboxylic acids and a process for producing same
- L28 ANSWER 19 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN DUPLICATE 1  
TI Method of manufacturing a magnetic storage medium
- L28 ANSWER 20 OF 69 USPATFULL on STN  
TI Liquid phase fluorination
- L28 ANSWER 21 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Synthesis of 1,1-dihydroperfluoroalkan-1-ols and their reaction with terephthaloyl chloride
- L28 ANSWER 22 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
TI The solid-like state of a confined liquid lubricant: deformation and time effects
- L28 ANSWER 23 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
TI The effect of adhesion on the rheological and frictional behavior of a confined lubricant film
- L28 ANSWER 24 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Manufacture of amides and esters of perfluoropolyoxyalkylenesulfonic or -carboxylic acids
- L28 ANSWER 25 OF 69 USPATFULL on STN  
TI Liquid phase fluorination
- L28 ANSWER 26 OF 69 USPATFULL on STN  
TI Curing fluorocarbon elastomers
- L28 ANSWER 27 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Cocyclotrimerization of mono- and dinitriles of perfluorocarboxylic acids under high pressure
- L28 ANSWER 28 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Development of quantitative structure-activity relationships for perfluoropolyalkyl ethers
- L28 ANSWER 29 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN DUPLICATE 2  
TI Liquid-phase fluorination
- L28 ANSWER 30 OF 69 USPATFULL on STN  
TI Liquid-phase fluorination

10/630,698

- L28 ANSWER 31 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN DUPLICATE 3  
TI Features of cyclotrimerization of perfluoroalkyl- and perfluorooxaalkylacetylenes
- L28 ANSWER 32 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
TI From static to kinetic friction in confined liquid films
- L28 ANSWER 33 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
TI New fluorinated oligomers and polymers based on (perfluoroalkyl)- and (perfluorooxyalkylene)acetylenes
- L28 ANSWER 34 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Lubricants for magnetic recording medium
- L28 ANSWER 35 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Curable fluororubber compositions containing fluorinated polyethers
- L28 ANSWER 36 OF 69 USPATFULL on STN  
TI Perfluorinated polyethers and process for their preparation
- L28 ANSWER 37 OF 69 USPATFULL on STN  
TI Liquid phase fluorination
- L28 ANSWER 38 OF 69 USPATFULL on STN  
TI Novel perfluorinated polyethers and process for their preparation
- L28 ANSWER 39 OF 69 USPATFULL on STN  
TI Novel perfluorinated polyethers and process for their preparation
- L28 ANSWER 40 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Colloid chemical properties of aqueous solutions of derivatives of perfluorooxaalkylcarboxylic acids based on oligomers of tetrafluoroethylene oxide
- L28 ANSWER 41 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Perfluoro tertiary alcohols. I. Synthesis of high molecular weight perfluorinated monoketones and tertiary alcohols
- L28 ANSWER 42 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Preparation of perfluoroacetal and perfluoroketal compounds and use thereof in thermal shock testing
- L28 ANSWER 43 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Perfluorination of hydrogen-containing compounds
- L28 ANSWER 44 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Perfluorination of alcohol ethoxylates
- L28 ANSWER 45 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN DUPLICATE 4  
TI Preparation of carbonyl fluoride compounds
- L28 ANSWER 46 OF 69 USPATFULL on STN  
TI Perfluoro-keto-ylids as precursors of polychloroketones, 1,2-diketones and quinoxalines
- L28 ANSWER 47 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN DUPLICATE 5  
TI Fluoroacrylate polymers and copolymers for manufacture of contact lenses.
- L28 ANSWER 48 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Lubricating oils for refrigerating compressors
- L28 ANSWER 49 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN

10/630,698

- TI Manufacture of magnetic memory disks
- L28 ANSWER 50 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Optical pickup activator
- L28 ANSWER 51 OF 69 USPATFULL on STN  
TI Perfluoro-keto-ylids as precursors of polychloroketones, 1,2-diketones and quinoxalines
- L28 ANSWER 52 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN DUPLICATE 6  
TI Perfluoro-keto-ylids as precursors of polychloroketones, 1,2-diketones and quinoxalines
- L28 ANSWER 53 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN DUPLICATE 7  
TI Perfluorocarbon ethers from a high-molecular-weight polyether
- L28 ANSWER 54 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Magnetic recording medium having a perfluoropolyether polymer protective coating
- L28 ANSWER 55 OF 69 USPATFULL on STN  
TI Magnetic recording medium having a perfluoropolyether polymer protective coating
- L28 ANSWER 56 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Fluoroalkylene ether silicate/viton GLT blends: an approach toward improved low temperature flexibility
- L28 ANSWER 57 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Hydrolytically stable fluorocarbon ether bibenzoxazole polymers
- L28 ANSWER 58 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
TI F-Phenylalkylene oxide diacetylenes
- L28 ANSWER 59 OF 69 USPATFULL on STN  
TI F-Phenylalkylene oxide diacetylenes
- L28 ANSWER 60 OF 69 USPATFULL on STN  
TI Fluoroalkyleneether silicate copolymers
- L28 ANSWER 61 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Fluoroalkyleneether silicate copolymers
- L28 ANSWER 62 OF 69 USPATFULL on STN  
TI Hybrid perfluoroalkylene ether thioimidate ester monomers
- L28 ANSWER 63 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Synthesis of the perfluoropoly(ethylene glycol) ethers by direct fluorination
- L28 ANSWER 64 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Hybrid perfluoroalkylene ether thioimidate ester monomers
- L28 ANSWER 65 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Synthesis of perfluoropoly(ethylene glycol) ethers CF<sub>3</sub>[OCF<sub>2</sub>CF<sub>2</sub>]<sub>n</sub>ORf (Rf = CF<sub>3</sub> or C<sub>2</sub>F<sub>5</sub>; n = 1-5)
- L28 ANSWER 66 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
TI The addition of tetrafluoroethylene oxide to F-glutaryl fluoride. Relative reactivities of acid fluorides
- L28 ANSWER 67 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN

10/630,698

TI Nature of the addition of perfluoroolefin oxides to perfluorodicarboxylic acid difluorides

L28 ANSWER 68 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Oil repellent polyfluoropolyoxo-alkyl phosphates

L28 ANSWER 69 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
TI Fluorocarbon ethers of tetrafluoroethylene epoxide

=> 2,16,17,20,25,29,30,36,37,38,39,44,45,53,65,66,67 bib ab fhitstr  
2,16,17,20,25,29,30,36,37,38,39,44,45,53,65,66,67 IS NOT A RECOGNIZED COMMAND  
The previous command name entered was not recognized by the system.  
For a list of commands available to you in the current file, enter  
"HELP COMMANDS" at an arrow prompt (=>).

=> d 2,16,17,20,25,29,30,36,37,38,39,44,45,53,65,66,67 bib ab fhitstr

L28 ANSWER 2 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN

AN 2004:75006 CAPLUS

DN 140:375766

TI Perfluoropoly-ether/peroxide compounds: spectroscopic studies and quantum chemical calculations

AU Radice, S.; Toniolo, P.; Barchiesi, E.; Guarda, P. A.; Tommasini, M.; Castiglioni, C.

CS Solvay Solexis, Bollate (MI), 20021, Italy

SO Journal of Fluorine Chemistry (2004), 125(2), 151-164

CODEN: JFLCAR; ISSN: 0022-1139

PB Elsevier Science B.V.

DT Journal

LA English

AB Perfluoropolyethers (PFPEs) are a class of high performance materials used in a wide range of applications (refrigeration, lubrication, semiconductor industry, etc.). PFPEs containing peroxidic units are intermediate materials for the preparation of com. end products. In this work we study the spectroscopic properties of ether and peroxides linkages in this class of compds.; NMR (NMR) spectra are discussed, FT-Raman data presented. Quantum chemical calcns. on model mols. were used as a tool for the interpretation of the Raman exptl. data and phys.-chemical properties.

IT 67584-24-1

RL: PRP (Properties)

(model compound; spectroscopic studies and quantum chemical calcns.  
perfluoropolyether/peroxide compds. prepared by oxidative polymerization of tetrafluoroethylene)

RN 67584-24-1 CAPLUS

CN 2,5,8,11,14,17,20-Heptaoxaheneicosane, 1,1,1,3,3,4,4,6,6,7,7,9,9,10,10,12,  
12,13,13,15,15,16,16,18,18,19,19,21,21,21-triacontafluoro- (9CI) (CA  
INDEX NAME)

PAGE 1-A

F<sub>3</sub>C—O—CF<sub>2</sub>—CF<sub>2</sub>—O—CF<sub>2</sub>—CF<sub>2</sub>—O—CF<sub>2</sub>—CF<sub>2</sub>—O—CF<sub>2</sub>—CF<sub>2</sub>—

PAGE 1-B

—O—CF<sub>2</sub> CF<sub>2</sub> O—CF<sub>3</sub>

RE.CNT 26 THERE ARE 26 CITED REFERENCES AVAILABLE FOR THIS RECORD

10/630,698

ALL CITATIONS AVAILABLE IN THE RE FORMAT

L28 ANSWER 16 OF 69 USPATFULL on STN  
AN 1998:55015 USPATFULL  
TI Liquid-phase fluorination  
IN Bierschenk, Thomas R., Round Rock, TX, United States  
Juhlke, Timothy J., Round Rock, TX, United States  
Kawa, Hajimu, Austin, TX, United States  
Lagow, Richard J., Austin, TX, United States  
PA Exfluor Research Corporation, Round Rock, TX, United States (U.S.  
corporation)  
PI US 5753776 19980519  
AI US 1995-471031 19950606 (8)  
RLI Continuation of Ser. No. US 1994-258708, filed on 13 Jun 1994, now  
patented, Pat. No. US 5461117, issued on 24 Oct 1995 which is a  
continuation of Ser. No. US 1993-28721, filed on 8 Mar 1993, now  
patented, Pat. No. US 5322904, issued on 21 Jun 1994 which is a  
continuation-in-part of Ser. No. US 1992-823837, filed on 17 Jan 1992,  
now abandoned which is a continuation of Ser. No. US 1989-414119, filed  
on 28 Sep 1989, now patented, Pat. No. US 5093432, issued on 3 Mar 1992  
which is a continuation-in-part of Ser. No. US 1988-250376, filed on 28  
Sep 1988, now abandoned  
DT Utility  
FS Granted  
EXNAM Primary Examiner: Krass, Frederick  
LREP Hamilton, Brook, Smith & Reynolds, P.C.  
CLMN Number of Claims: 35  
ECL Exemplary Claim: 1  
DRWN 2 Drawing Figure(s); 2 Drawing Page(s)  
LN.CNT 2151  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.  
AB This invention pertains to a method for liquid phase fluorination for  
perfluorination of a wide variety of hydrogen-containing compounds.  
IT 130085-03-9P  
(preparation of)  
RN 130085-03-9 USPATFULL  
CN 5,8,11,13,16,19-Hexaoxatricosane, 1,1,1,2,2,3,3,4,4,6,6,7,7,9,9,10,10,12,1  
2,14,14,15,15,17,17,18,18,20,20,21,21,22,22,23,23,23-hexatriacontafluoro-  
(9CI) (CA INDEX NAME)

PAGE 1-A

F<sub>3</sub>C—(CF<sub>2</sub>)<sub>3</sub>—O—CF<sub>2</sub>—CF<sub>2</sub> O—CF<sub>2</sub>—CF<sub>2</sub>—O—CF<sub>2</sub>—O—CF<sub>2</sub> CF<sub>2</sub> O—CF<sub>2</sub>—

PAGE 1-B

—CF<sub>2</sub>—O—(CF<sub>2</sub>)<sub>3</sub>—CF<sub>3</sub>

L28 ANSWER 17 OF 69 USPATFULL on STN  
AN 97:91604 USPATFULL  
TI Liquid-phase fluorination  
IN Bierschenk, Thomas R., Round Rock, TX, United States  
Juhlke, Timothy, Round Rock, TX, United States  
Kawa, Hajimu, Austin, TX, United States  
Lagow, Richard J., Austin, TX, United States  
PA Exfluor Research Corporation, Round Rock, TX, United States (U.S.  
corporation)

10/630,698

PI US 5674949 19971007  
AI US 1995-466798 19950606 (8)  
RLI Continuation of Ser. No. US 1994-240225, filed on 10 May 1994, now patented, Pat. No. US 5571870, issued on 5 Nov 1996 which is a continuation of Ser. No. US 1992-823836, filed on 17 Jan 1992, now patented, Pat. No. US 5322903, issued on 21 Jun 1994 which is a continuation of Ser. No. US 1989-414119, filed on 28 Sep 1989, now patented, Pat. No. US 5093432, issued on 3 Mar 1992 which is a continuation-in-part of Ser. No. US 1988-250376, filed on 28 Sep 1988, now abandoned  
DT Utility  
FS Granted  
EXNAM Primary Examiner: Krass, Frederick  
LREP Hamilton, Brook, Smith & Reynolds, P.C.  
CLMN Number of Claims: 20  
ECL Exemplary Claim: 1  
DRWN 2 Drawing Figure(s); 2 Drawing Page(s)  
LN.CNT 2088  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.  
AB This invention pertains to a method for liquid-phase fluorination for perfluorination of a wide variety of hydrogen-containing compounds.  
IT 130085-03-9P  
    (preparation of)  
RN 130085-03-9 USPATFULL  
CN 5,8,11,13,16,19-Hexaoxatricosane, 1,1,1,2,2,3,3,4,4,6,6,7,7,9,9,10,10,12,1  
    2,14,14,15,15,17,17,18,18,20,20,21,21,22,22,23,23-hexatriacontafluoro-  
    (9CI) (CA INDEX NAME)

PAGE 1-A

F<sub>3</sub>C—(CF<sub>2</sub>)<sub>3</sub>—O—CF<sub>2</sub>—CF<sub>2</sub>—O—CF<sub>2</sub>—CF<sub>2</sub>—O—CF<sub>2</sub>—CF<sub>2</sub>—O—CF<sub>2</sub>—

PAGE 1-B

CF<sub>2</sub> O (CF<sub>2</sub>)<sub>3</sub>—CF<sub>3</sub>

L28 ANSWER 20 OF 69 USPATFULL on STN  
AN 96:101634 USPATFULL  
TI Liquid phase fluorination  
IN Bierschenk, Thomas R., Round Rock, TX, United States  
    Juhlke, Timothy, Round Rock, TX, United States  
    Kawa, Hajimu, Austin, TX, United States  
    Lagow, Richard J., Austin, TX, United States  
PA Exfluor Research Corporation, Round Rock, TX, United States (U.S.  
    corporation)  
PI US 5571870 19961105  
AI US 1994-240225 19940510 (8)  
DCD 20110621  
RLI Continuation of Ser. No. US 1992-823836, filed on 17 Jan 1992, now patented, Pat. No. US 5322903, issued on 21 Jun 1994 which is a continuation of Ser. No. US 1989-414119, filed on 28 Sep 1989, now patented, Pat. No. US 5093432, issued on 3 Mar 1992 which is a continuation-in-part of Ser. No. US 1988-250376, filed on 28 Sep 1988, now abandoned  
DT Utility  
FS Granted  
EXNAM Primary Examiner: Krass, Frederick

10/630,698

LREP Hamilton, Brook, Smith & Reynolds, P.C.

CLMN Number of Claims: 28

ECL Exemplary Claim: 1

DRWN 2 Drawing Figure(s); 2 Drawing Page(s)

LN.CNT 2065

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB This invention pertains to a method for liquid phase fluorination for perfluorination of a wide variety of hydrogen-containing compounds.

IT 130085-03-9P

(preparation of)

RN 130085-03-9 USPATFULL

CN 5,8,11,13,16,19-Hexaoxatricosane, 1,1,1,2,2,3,3,4,4,6,6,7,7,9,9,10,10,12,1  
2,14,14,15,15,17,17,18,18,20,20,21,21,22,22,23,23-hexatriacontafluoro-  
(9CI) (CA INDEX NAME)

PAGE 1-A

F<sub>3</sub>C—(CF<sub>2</sub>)<sub>3</sub>—O—CF<sub>2</sub>—CF<sub>2</sub>—O—CF<sub>2</sub>—CF<sub>2</sub>—O—CF<sub>2</sub>—O—CF<sub>2</sub>—CF<sub>2</sub>—O—CF<sub>2</sub>—

PAGE 1-B

— CF<sub>2</sub>—O—(CF<sub>2</sub>)<sub>3</sub>—CF<sub>3</sub>

L28 ANSWER 25 OF 69 USPATFULL on STN

AN 95:94983 USPATFULL

TI Liquid phase fluorination

IN Bierschenk, Thomas R., Round Rock, TX, United States

Juhlke, Timothy J., Round Rock, TX, United States

Kawa, Hajimu, Austin, TX, United States

Lagow, Richard J., Austin, TX, United States

PA Exfluor Research Corporation, Austin, TX, United States (U.S.  
corporation)

PI US 5461117 19951024

AI US 1994-258708 19940613 (8)

RLI Continuation of Ser. No. US 1993-28721, filed on 8 Mar 1993, now  
patented, Pat. No. US 5322904 which is a continuation-in-part of Ser.  
No. US 1992-823837, filed on 17 Jan 1992, now abandoned which is a  
continuation of Ser. No. US 1989-414119, filed on 28 Sep 1989, now  
patented, Pat. No. US 5094432, issued on 3 Mar 1992 which is a  
continuation-in-part of Ser. No. US 1988-250376, filed on 28 Sep 1988,  
now abandoned

DT Utility

FS Granted

EXNAM Primary Examiner: Krass, Frederick

LREP Hamilton, Brook, Smith & Reynolds

CLMN Number of Claims: 30

ECL Exemplary Claim: 1

DRWN 2 Drawing Figure(s); 2 Drawing Page(s)

LN.CNT 2106

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB This invention pertains to a method for liquid phase fluorination for perfluorination of a wide variety of hydrogen-containing compounds.

IT 130085-03-9P

(preparation of)

RN 130085-03-9 USPATFULL

CN 5,8,11,13,16,19-Hexaoxatricosane, 1,1,1,2,2,3,3,4,4,6,6,7,7,9,9,10,10,12,1  
2,14,14,15,15,17,17,18,18,20,20,21,21,22,22,23,23-hexatriacontafluoro-

10/630,698

(9CI) (CA INDEX NAME)

PAGE 1-A

F<sub>3</sub>C—(CF<sub>2</sub>)<sub>3</sub>—O—CF<sub>2</sub>—CF<sub>2</sub>—O—CF<sub>2</sub>—CF<sub>2</sub>—O—CF<sub>2</sub>—CF<sub>2</sub>—O—CF<sub>2</sub>—

PAGE 1-B

— CF<sub>2</sub>—O—(CF<sub>2</sub>)<sub>3</sub>—CF<sub>3</sub>

L28 ANSWER 29 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN DUPLICATE 2

AN 1994:701682 CAPLUS

DN 121:301682

TI Liquid-phase fluorination

IN Bierschenk, Thomas R.; Juhlke, Timothy; Kawa, Hajimu; Lagow, Richard J.

PA Exfluor Research Corp., USA

SO U.S., 24 pp. Cont.-in-part of U.S. Ser. No. 822,637, abandoned.

CODEN: USXXAM

DT Patent

LA English

FAN.CNT 3

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 5332790	A	19940726	US 1993-28682	19930308
	US 5093432	A	19920303	US 1989-414119	19890928
	US 5322903	A	19940621	US 1992-823836	19920117
	US 5571870	A	19961105	US 1994-240225	19940510
	US 5674949	A	19971007	US 1995-466798	19950606

PRAI US 1988-250376 19880928  
US 1989-414119 19890928  
US 1992-822637 19920117  
US 1992-823836 19920117  
US 1994-240225 19940510

AB A method for replacing essentially all H atoms of H-containing compds. with F atoms comprises (a) continuously introducing a H-containing compound into a liquid perfluorocarbon, perhalogenated chlorofluorocarbon or chloro fluoro ether medium while agitating the medium so that the H-containing compound is dissolved or dispersed within the liquid medium; (b) introducing F gas diluted with an inert gas into the liquid medium without illumination with UV light to establish fluorination conditions wherein the liquid medium and F in the vapor space above the liquid medium do not form a flammable mixture; (c) continuing the introduction of F gas diluted with an inert gas until essentially all of the H atoms of the H-containing compound have been replaced with F atoms without substantial oligomerization or polymerization of the H-containing compound. Perfluorinated acids (such as C<sub>7</sub>F<sub>15</sub>CO<sub>2</sub>H), perfluorinated polyethylene glycol and polypropylene glycol and their derivs. were prepared

IT 130085-03-9P

RL: IMF (Industrial manufacture); PREP (Preparation)  
(liquid-phase perfluorination)

RN 130085-03-9 CAPLUS

CN 5,8,11,13,16,19-Hexaoxatricosane, 1,1,1,2,2,3,3,4,4,6,6,7,7,9,9,10,10,12,12,14,14,15,15,17,17,18,18,20,20,21,21,22,22,23,23-hexatriacontafluoro-(9CI) (CA INDEX NAME)

10/630,698

PAGE 1-A

F<sub>3</sub>C—(CF<sub>2</sub>)<sub>3</sub>—O—CF<sub>2</sub>—CF<sub>2</sub>—O—CF<sub>2</sub>—CF<sub>2</sub>—O—CF<sub>2</sub>—O—CF<sub>2</sub>—CF<sub>2</sub>—O—CF<sub>2</sub>—

PAGE 1-B

—CF<sub>2</sub>—O—(CF<sub>2</sub>)<sub>3</sub>—CF<sub>3</sub>

L28 ANSWER 30 OF 69 USPATFULL on STN  
AN 94:53503 USPATFULL  
TI Liquid-phase fluorination  
IN Bierschenk, Thomas R., Round Rock, TX, United States  
Juhlke, Timothy, Round Rock, TX, United States  
Kawa, Hajimu, Austin, TX, United States  
Lagow, Richard J., Austin, TX, United States  
PA Exfluor Research Corporation, Austin, TX, United States (U.S.  
corporation)  
PI US 5322903 19940621  
AI US 1992-823836 19920117 (7)  
RLI Continuation of Ser. No. US 1989-414119, filed on 28 Sep 1989, now  
patented, Pat. No. US 5093432, issued on 3 Mar 1992 which is a  
continuation-in-part of Ser. No. US 1988-250376, filed on 28 Sep 1988,  
now abandoned  
DT Utility  
FS Granted  
EXNAM Primary Examiner: Krass, Frederick  
LREP Hamilton, Brook, Smith & Reynolds  
CLMN Number of Claims: 20  
ECL Exemplary Claim: 1  
DRWN 2 Drawing Figure(s); 2 Drawing Page(s)  
LN.CNT 1950  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.  
AB This invention pertains to a method for liquid phase fluorination for  
perfluorination of a wide variety of hydrogen-containing compounds.  
IT 130085-03-9P  
(preparation of)  
RN 130085-03-9 USPATFULL  
CN 5,8,11,13,16,19-Hexaoxatricosane, 1,1,1,2,2,3,3,4,4,6,6,7,7,9,9,10,10,12,1  
2,14,14,15,15,17,17,18,18,20,20,21,21,22,22,23,23,23-hexatriacontafluoro-  
(9CI) (CA INDEX NAME)

PAGE 1-A

F<sub>3</sub>C—(CF<sub>2</sub>)<sub>3</sub>—O—CF<sub>2</sub>—CF<sub>2</sub>—O—CF<sub>2</sub>—CF<sub>2</sub>—O—CF<sub>2</sub>—O—CF<sub>2</sub>—CF<sub>2</sub>—O—CF<sub>2</sub>—

PAGE 1-B

—CF<sub>2</sub>—O—(CF<sub>2</sub>)<sub>3</sub>—CF<sub>3</sub>

L28 ANSWER 36 OF 69 USPATFULL on STN  
AN 92:46791 USPATFULL  
TI Perfluorinated polyethers and process for their preparation  
IN Kalota, Dennis J., Fenton, MO, United States

10/630,698

McConaghy, Jr., John S., St. Louis, MO, United States  
Foerst, Paul W., Chesterfield, MO, United States  
Liu, Paul H., Chesterfield, MO, United States  
Feher, Jr., Frank R., Belleville, IL, United States  
PA Monsanto Company, St. Louis, MO, United States (U.S. corporation)  
PI US 5120459 19920609  
AI US 1990-498055 19900323 (7)  
RLI Division of Ser. No. US 1989-150963, filed on 29 Jan 1989, now abandoned  
DT Utility  
FS Granted  
EXNAM Primary Examiner: McAvoy, Ellen  
LREP Brooks, W. W.  
CLMN Number of Claims: 2  
ECL Exemplary Claim: 1  
DRWN 2 Drawing Figure(s); 2 Drawing Page(s)  
LN.CNT 535  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.  
AB Perfluorinated polyethers having the formula

R.sub.f 0--(CF.sub.2 CF.sub.2 0).sub.n --R'.sub.f

wherein n is an integer of 1-11 and each of R.sub.f and R'.sub.f is a perfluorinated C.sub.1 -C.sub.5 alkyl radical, dimers of such polyethers and carbon to carbon intramolecularly coupled cyclic derivatives of such polyethers are produced by direct fluorination of the polyethers in an inert solvent. Compositions of the perfluorinated polyethers and their derivatives are useful as functional fluids.

IT 125662-66-0P  
(manufacture of, solvent for)  
RN 125662-66-0 USPATFULL  
CN 2,5,8,11,14,17,20,23-Octaoxatetracosane, 1,1,1,3,3,4,4,6,6,7,7,9,9,10,10,11  
2,12,13,13,15,15,16,16,18,18,19,19,21,21,22,22,24,24,24-  
tetratriacontafluoro- (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 1 - B

$$\cdots \text{O} - \text{CF}_2 - \text{CF}_2 - \text{O} - \text{CF}_2 - \text{CF}_2 - \text{O} - \text{CF}_3$$

L28 ANSWER 37 OF 69 USPATFULL on STN  
AN 92:17225 USPATFULL  
TI Liquid phase fluorination  
IN Bierschenk, Thomas R., Round Rock, TX, United States  
      Juhlke, Timothy, Round Rock, TX, United States  
      Kawa, Hajimu, Austin, TX, United States  
      Lagow, Richard J., Austin, TX, United States  
PA Exfluor Research Corporation, Austin, TX, United States (U.S.  
      corporation)  
PI US 5093432                   19920303  
AI US 1989-414119               19890928 (7)  
RLI Continuation-in-part of Ser. No. US 1988-250376, filed on 28 Sep 1988,  
      now abandoned  
DT Utility  
FS Granted

10/630,698

EXNAM Primary Examiner: Kight, III, John; Assistant Examiner: Krass, Frederick  
LREP Hamilton, Brook, Smith & Reynolds  
CLMN Number of Claims: 21  
ECL Exemplary Claim: 1  
DRWN 2 Drawing Figure(s); 2 Drawing Page(s)  
LN.CNT 2057

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB This invention pertains to a method for liquid phase fluorination for perfluorination of a wide variety of hydrogen-containing compounds.

IT 130085-03-9P

(preparation of)

RN 130085-03-9 USPATFULL

CN 5,8,11,13,16,19-Hexaoxatricosane, 1,1,1,2,2,3,3,4,4,6,6,7,7,9,9,10,10,12,1  
2,14,14,15,15,17,17,18,18,20,20,21,21,22,22,23,23-hexatriacontafluoro-  
(9CI) (CA INDEX NAME)

PAGE 1-A

F<sub>3</sub>C—(CF<sub>2</sub>)<sub>3</sub>—O—CF<sub>2</sub>—CF<sub>2</sub>—O—CF<sub>2</sub>—CF<sub>2</sub>—O—CF<sub>2</sub>—CF<sub>2</sub>—O—CF<sub>2</sub>—

PAGE 1-B

—CF<sub>2</sub> O (CF<sub>2</sub>)<sub>3</sub>—CF<sub>3</sub>

L28 ANSWER 38 OF 69 USPATFULL on STN  
AN 91:106001 USPATFULL  
TI Novel perfluorinated polyethers and process for their preparation  
IN Kalota, Dennis J., Fenton, MO, United States  
McConaghy, Jr., John S., St. Louis, MO, United States  
Foerst, Paul W., Chesterfield, MO, United States  
Liu, Paul H., Chesterfield, MO, United States  
Feher, Jr., Frank R., Belleville, IL, United States  
PA Monsanto Company, St. Louis, MO, United States (U.S. corporation)  
PI US 5076949 19911231  
AI US 1990-498124 19900323 (7)  
RLI Division of Ser. No. US 1989-150963, filed on 29 Jan 1989, now abandoned  
DT Utility  
FS Granted  
EXNAM Primary Examiner: Willis, Jr., Prince; Assistant Examiner: McAvoy, Ellen  
LREP Brooks, W. W.  
CLMN Number of Claims: 5  
ECL Exemplary Claim: 1  
DRWN 2 Drawing Figure(s); 2 Drawing Page(s)  
LN.CNT 549  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.  
AB Perfluorinated polyethers having the formula

R.<sub>sub.f</sub> O--(CF.<sub>sub.2</sub> CF.<sub>sub.2</sub> O).<sub>sub.n</sub>--R'.<sub>sub.f</sub>

wherein n is an integer of 1-11 and each of R.<sub>sub.f</sub> and R'.<sub>sub.f</sub> is a perfluorinated C.<sub>sub.1</sub>-C.<sub>sub.5</sub>-alkyl radical, dimers of such polyethers and carbon to carbon intramolecularly coupled cyclic derivatives of such polyethers are produced by direct fluorination of the polyethers in an inert solvent. Compositions of the perfluorinated polyethers and their derivatives are useful as functional fluids.

IT 125662-66-0P

(manufacture of, solvent for)

10/630,698

RN 125662-66-0 USPATFULL  
CN 2,5,8,11,14,17,20,23-Octaoxatetracosane, 1,1,1,3,3,4,4,6,6,7,7,9,9,10,10,1  
2,12,13,13,15,15,16,16,18,18,19,19,21,21,22,22,24,24,24-  
tetra(triaconta)fluoro- (9CI) (CA INDEX NAME)

PAGE 1-A

F<sub>3</sub>C—O—CF<sub>2</sub>—CF<sub>2</sub>—O—CF<sub>2</sub>—CF<sub>2</sub>—O—CF<sub>2</sub>—CF<sub>2</sub>—O—CF<sub>2</sub>—CF<sub>2</sub>—O—CF<sub>2</sub>—CF<sub>2</sub>—

PAGE 1-B

—O—CF<sub>2</sub>—CF<sub>2</sub>—O—CF<sub>2</sub>—CF<sub>2</sub>—O—CF<sub>3</sub>

L28 ANSWER 39 OF 69 USPATFULL on STN  
AN 91:17276 USPATFULL  
TI Novel perfluorinated polyethers and process for their preparation  
IN Kalota, Dennis J., Fenton, MO, United States  
McConaghy, Jr., John S., St. Louis, MO, United States  
Foerst, Paul W., Chesterfield, MO, United States  
Liu, Paul H., Chesterfield, MO, United States  
Feher, Jr., Frank R., Belleville, IL, United States  
PA Monsanto Company, St. Louis, MO, United States (U.S. corporation)  
PI US 4996369 19910226  
AI US 1990-498057 19900523 (7)  
RLI Division of Ser. No. US 1989-150963, filed on 29 Jan 1989  
DT Utility  
FS Granted  
EXNAM Primary Examiner: Mars, Howard T.  
LREP Brooks, W.  
CLMN Number of Claims: 1  
ECL Exemplary Claim: 1  
DRWN 2 Drawing Figure(s); 2 Drawing Page(s)  
LN.CNT 533  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.  
AB Perfluorinated polyethers having the formula

R.<sub>sub.f</sub> O--(CF.<sub>sub.2</sub> CF.<sub>sub.2</sub> O).<sub>sub.n</sub>--R'.<sub>sub.f</sub>

wherein n is an integer of 1-11 and each of R.<sub>sub.f</sub> and R'.<sub>sub.f</sub> is a perfluorinated C.<sub>sub.1</sub>-C.<sub>sub.5</sub>-alkyl radical, dimers of such polyethers and carbon to carbon intramolecularly coupled cyclic derivatives of such polyethers are produced by direct fluorination of the polyethers in an inert solvent. Compositions of the perfluorinated polyethers and their derivatives are useful as functional fluids.

IT 125662-66-0P  
(manufacture of, solvent for)  
RN 125662-66-0 USPATFULL  
CN 2,5,8,11,14,17,20,23-Octaoxatetracosane, 1,1,1,3,3,4,4,6,6,7,7,9,9,10,10,1  
2,12,13,13,15,15,16,16,18,18,19,19,21,21,22,22,24,24,24-  
tetra(triaconta)fluoro- (9CI) (CA INDEX NAME)

PAGE 1-A

F<sub>3</sub>C—O—CF<sub>2</sub>—CF<sub>2</sub>—O—CF<sub>2</sub>—CF<sub>2</sub>—O—CF<sub>2</sub>—CF<sub>2</sub>—O—CF<sub>2</sub>—CF<sub>2</sub>—

- O-CF<sub>2</sub>-CF<sub>2</sub>-O-CF<sub>2</sub>-CF<sub>2</sub>-O-CF<sub>3</sub>

L28 ANSWER 44 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
 AN 1990:121089 CAPLUS  
 DN 112:121089  
 TI Perfluorination of alcohol ethoxylates  
 IN Feher, Frank Ronald; Foerst, Paul Wayne; Liu, Paul Ho; Kalota, Dennis  
 Jerome; McConaghay, John Stead, Jr.  
 PA Monsanto Co., USA  
 SO Eur. Pat. Appl., 12 pp.  
 CODEN: EPXXDW  
 DT Patent  
 LA English  
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	EP 332601 R: AT, BE, CH, DE, ES, FR, GB, GR, IT, LI, LU, NL, SE AU 8928869 AU 607579 JP 01225628 US 5076949 US 5120459 US 4996369 JP 06025404 JP 07086139 JP 06025405 JP 07086140 JP 06080773 JP 07088423	A1 A1 B2 A2 A A A A2 B4 A2 B4 A2 B4 A2 B4	19890913 19890803 19910307 19890908 19911231 19920609 19910226 19940201 19950920 19940201 19950920 19940322 19950927	EP 1989-870017 AU 1989-28869 JP 1989-19415 US 1990-498124 US 1990-498055 US 1990-498057 JP 1993-115041 JP 1993-115042 JP 1993-115043	19890127 19890127 19890127 19900323 19900323 19900523 19930517 19930517 19930517 19930517

PRAI US 1988-150963 19880129

AB The title compds. RO(CF<sub>2</sub>CF<sub>2</sub>O)<sub>n</sub>R<sub>1</sub> (R, R<sub>1</sub> = perfluorinated C<sub>1</sub>-5 alkyl; n = 1-11) are prepared by reacting F(g) with alc. ethoxylates R<sub>2</sub>O(CH<sub>2</sub>CH<sub>2</sub>O)<sub>n</sub>R<sub>3</sub> (R<sub>2</sub>, R<sub>3</sub> = C<sub>1</sub>-5 alkyl; n = 1-11) in an inert solvent and separating the product. A process schematic and a reactor diagram are presented. Thus, 250 g heptaglyme and a slurry of 1110 g NaF in 4 L 1,1,2-trichloro-1,2,2-trifluoroethane was charged into a stirred (1200 rpm) reactor, a mixture of F and N added at 15-25° for 4 h, 7105 g of the fluorinated oil intermediate (5-10% H content) was charged into a reactor with 300 g NaF, the oil reacted with F at 31-128° for 177 min, the treated oil reacted with F at 29-253° for 230 min, and distilled to give a title product having average mol. weight 1000, b.p. (760 torr) 215°, pour point -25°, and d<sub>20</sub> 1.72 g/mL. The distillation bottoms contained perfluoroheptaglyme dimer and oligomers having average mol. weight 1900, b.p. 200°/4 torr, pour point -70°, and d<sub>20</sub> 1.81 g/mL.

IT 125662-66-0P

RL: IMF (Industrial manufacture); PREP (Preparation)  
 (manufacture of, solvent for)

RN 125662-66-0 CAPLUS

CN 2,5,8,11,14,17,20,23-Octaoxatetracosane, 1,1,1,3,3,4,4,6,6,7,7,9,9,10,10,1  
 2,12,13,13,15,15,16,16,18,18,19,19,21,21,22,22,24,24-tetratriacontafluoro- (9CI) (CA INDEX NAME)

PAGE 1-A

$$\text{F}_3\text{C}-\text{O}-\text{CF}_2-\text{CF}_2-\text{O}-\text{CF}_2-\text{CF}_2-\text{O}-\text{CF}_2-\text{CF}_2-\text{O}-\text{CF}_2-\text{CF}_2-\text{O}-\text{CF}_2-\text{CF}_2-$$

PAGE 1-B

$$-\text{O}-\text{CF}_2-\text{CF}_2-\text{O}-\text{CF}_2-\text{CF}_2-\text{O}-\text{CF}_3$$

L28 ANSWER 45 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN DUPLICATE 4

AN 1989:438876 CAPLUS

DN 111:38876

TI Preparation of carbonyl fluoride compounds

IN Okabe, Jun; Tatsu, Haruyoshi

PA Nippon Mectron Co., Ltd., Japan

SO U.S., 7 pp.

CODEN: USXXAM

DT Patent

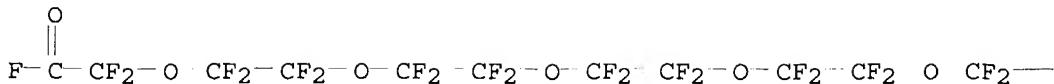
LA English

FAN.CNT 1

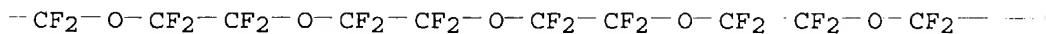
	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 4769184	A	19880906	US 1987-121135	19871116
	JP 01066139	A2	19890313	JP 1987-222946	19870908
	JP 08019035	B4	19960228		
	JP 01093557	A2	19890412	JP 1987-249588	19871002
	JP 2726824	B2	19980311		
PRAI	JP 1987-222946		19870908		
	JP 1987-249588		19871002		
OS	MARPAT 111:38876				
AB	A process for producing XCOF (I; X = F, CF <sub>3</sub> ) or I (X = CF <sub>3</sub> CF <sub>2</sub> ), useful as intermediates for producing perfluoro(alkyl vinyl ethers) which are monomers for producing F-containing resins, F-containing rubber, etc., comprised				
	thermally decomposing RfO(CF <sub>2</sub> CF <sub>2</sub> O) <sub>a</sub> (CF <sub>2</sub> O) <sub>b</sub> (O) <sub>c</sub> Rf' (Rf = perfluoroalkyl; Rf' = COF, CF <sub>3</sub> ; the CF <sub>2</sub> O and O groups are distributed at random; a, b ≠ 0; c can be 0; a + b + c ≤ .apprx.200) or RfO(CFXCF <sub>2</sub> O) <sub>n</sub> CFX'Y (X' = CF <sub>3</sub> , F, H; Y = COF, CO <sub>2</sub> H, CO <sub>2</sub> R, CF <sub>3</sub> ; R = alkyl; n = 1-50), resp. F <sub>2</sub> C:CF <sub>2</sub> and O <sub>2</sub> were irradiated with UV to give F <sub>3</sub> CO(CF <sub>2</sub> OF <sub>2</sub> O) <sub>8</sub> (CF <sub>2</sub> O) <sub>24</sub> O <sub>2</sub> .4COF, thermal decomposition of which at 200° over activated C gave a mixture of 78.2% COF <sub>2</sub> and 21.8% F <sub>3</sub> CCOF. I (X = F, CF <sub>3</sub> ) so produced contain no Cl-based impurities.				
IT	119214-96-9P				
	RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)				
	(preparation and reaction of, in synthesis of carbonyl fluorides)				
RN	119214-96-9	CAPLUS			
CN	3,6,9,12,15,18,21,24,27,30,33,36,39,42,45,48,51,54,57,60,63,66,69- Tricosaoxahenheptacontanoyl fluoride, 2,2,4,4,5,5,7,7,8,8,10,10,11,11,13,1 3,14,14,16,16,17,17,19,19,20,20,22,22,23,23,25,25,26,26,28,28,29,29,31,31, 32,32,34,34,35,35,37,37,38,38,40,40,41,41,43,43,44,44,46,46,47,47,49,49,50 ,50,52,52,53,53,55,55,56,56,58,58,59,59,61,61,62,62,64,64,65,65,67,67,68,6 8,70,70,71,71,71-pentanonacontafluoro- (9CI) (CA INDEX NAME)				

10/630,698

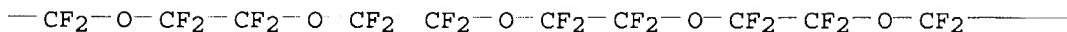
PAGE 1-A



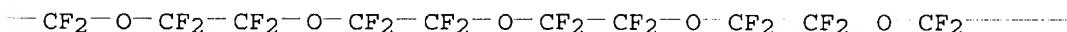
PAGE 1-B



PAGE 1-C



PAGE 1-D



PAGE 1-E



L28 ANSWER 53 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN DUPLICATE 7

AN 1986:5566 CAPLUS

DN 104:5566

TI Perfluorocarbon ethers from a high-molecular-weight polyether  
IN Lagow, Richard J.; Gerhardt, Glenn E.

PA University of Texas, USA

SO U.S., 16 pp. Cont. U.S. Ser. No. 139,181 abandoned.  
CODEN: USXXAM

DT Patent

LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 4523039	A	19850611	US 1983-563013	19831219
PRAI	US 1978-901905		19780501		
	US 1980-139181		19800411		

AB Fluorocarbon ethers were prepared by fluorination of a high mol. weight polyether with F<sub>2</sub> to produce a fluorinated polyether, which was depolymerd. by further treatment with F<sub>2</sub> at 55-210°. Thus, polyethylene exide

10/630,698

was fluoroinated with flowing F<sub>2</sub>-He, using LaMar techniques, at ambient temperature for 12 days, at 90° for 2 days, and at 110° for 7 days, to give a mixture of compds. including CF<sub>4</sub>, COF<sub>2</sub>, F<sub>3</sub>CO(CF<sub>2</sub>CF<sub>2</sub>O)<sub>n</sub>R (n = 1-6; R = CF<sub>3</sub>, C<sub>2</sub>F<sub>5</sub>) (all permutations), and C<sub>2</sub>F<sub>5</sub>O(CF<sub>2</sub>CF<sub>2</sub>O)<sub>m</sub>C<sub>2</sub>F<sub>5</sub> (m = 1-3), which were separated and characterized by IR, <sup>19</sup>F NMR, and mass spectroscopy.

IT 64028-08-6P

RL: SPN (Synthetic preparation); PREP (Preparation)  
(preparation of, by fluorination-depolymn. of polyether, and spectral characterization of)

RN 64028-08-6 CAPLUS

CN 2,5,8,11,14,17-Hexaoxaoctadecane, 1,1,1,2,2,4,4,6,6,7,7,9,9,10,10,12,12,13  
,13,15,15,16,16,18,18,18-hexacosafuoro- (9CI) (CA INDEX NAME)

PAGE 1-A

F<sub>3</sub>C—O—CF<sub>2</sub>—CF<sub>2</sub>—O—CF<sub>2</sub>—CF<sub>2</sub>—O—CF<sub>2</sub>—CF<sub>2</sub>—O—CF<sub>2</sub>—CF<sub>2</sub>—O—CF<sub>2</sub>—CF<sub>2</sub>—

PAGE 1-B

--O—CF<sub>3</sub>

L28 ANSWER 65 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
AN 1977:517527 CAPLUS

DN 87:117527

TI Synthesis of perfluoropoly(ethylene glycol) ethers CF<sub>3</sub>[OCF<sub>2</sub>CF<sub>2</sub>]<sub>n</sub>ORf (Rf = CF<sub>3</sub> or C<sub>2</sub>F<sub>5</sub>; n = 1-5)

AU Gerhardt, Glenn E.; Lagow, Richard J.

CS Dep. Chem., Univ. Texas, Austin, TX, USA

SO Journal of the Chemical Society, Chemical Communications (1977), (8), 259-60

CODEN: JCCCAT; ISSN: 0022-4936

DT Journal

LA English

AB Finely ground (<120 mesh) poly(ethylene oxide) reacted with elemental F, under conditions carefully regulated to fragment and perfluorinate the polyether system, to give CF<sub>3</sub>[O(CF<sub>2</sub>)<sub>2</sub>]<sub>n</sub>OR (R = CF<sub>3</sub>, C<sub>2</sub>F<sub>5</sub>, n = 1-5).

IT 64028-08-6P

RL: SPN (Synthetic preparation); PREP (Preparation)  
(preparation of)

RN 64028-08-6 CAPLUS

CN 2,5,8,11,14,17-Hexaoxaoctadecane, 1,1,1,2,2,4,4,6,6,7,7,9,9,10,10,12,12,13  
,13,15,15,16,16,18,18,18-hexacosafuoro- (9CI) (CA INDEX NAME)

PAGE 1-A

F<sub>3</sub>C—O—CF<sub>2</sub>—CF<sub>2</sub>—O—CF<sub>2</sub>—CF<sub>2</sub>—O—CF<sub>2</sub>—CF<sub>2</sub>—O—CF<sub>2</sub>—CF<sub>2</sub>—O—CF<sub>2</sub>—CF<sub>2</sub>—

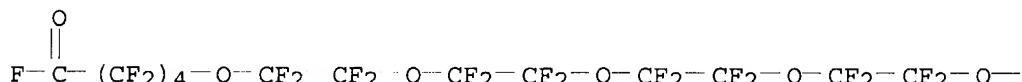
PAGE 1-B

--O—CF<sub>3</sub>

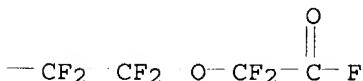
L28 ANSWER 66 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN

AN 1976:493374 CAPLUS  
 DN 85:93374  
 TI The addition of tetrafluoroethylene oxide to F-glutaryl fluoride.  
 Relative reactivities of acid fluorides  
 AU Anderson, R.; Baucom, K. B.; Psarras, T.; Snyder, C. E.; Cochoy, R. E.  
 CS PCR, Inc., Gainesville, FL, USA  
 SO Journal of Fluorine Chemistry (1976), 7(6), 581-8  
 CODEN: JFLCAR; ISSN: 0022-1139  
 DT Journal  
 LA English  
 AB Data obtained from the addition of tetrafluoroethylene oxide to F-glutaryl fluoride [FOC(CF<sub>2</sub>)<sub>3</sub>COF] show significant differences to exist between the relative reactivities of the acid fluoride groups involved. The order of reactivity is F-glutaryl fluoride > -OCF<sub>2</sub>COF > -CF<sub>2</sub>CF<sub>2</sub>COF.  
 IT 60127-05-1P  
 RL: SPN (Synthetic preparation); PREP (Preparation)  
 (preparation of)  
 RN 60127-05-1 CAPLUS  
 CN 3,6,9,12,15,18-Hexaoxatricosanediol difluoride,  
 2,2,4,4,5,5,7,7,8,8,10,10,11,11,13,13,14,14,16,16,17,17,19,19,20,20,21,21,  
 22,22-triacontafluoro- (9CI) (CA INDEX NAME)

PAGE 1-A



PAGE 1-B

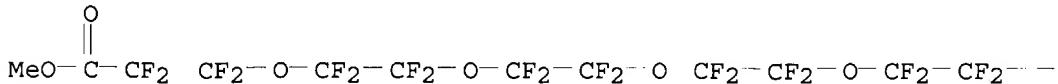


L28 ANSWER 67 OF 69 CAPLUS COPYRIGHT 2004 ACS on STN  
 AN 1974:26718 CAPLUS  
 DN 80:26718  
 TI Nature of the addition of perfluoroolefin oxides to perfluorodicarboxylic acid difluorides  
 AU Skoblikova, V. I.; Sass, V. P.; Ershov, A. E.; Senyushov, L. N.; Sokolov, L. F.; Berenblit, V. V.; Sokolov, S. V.  
 CS Vses. Nauchno-Issled Inst. Sint. Kauch., Leningrad, USSR  
 SO Zhurnal Organicheskoi Khimii (1973), 9(10), 2021-5  
 CODEN: ZORKAE; ISSN: 0514-7492  
 DT Journal  
 LA Russian  
 AB FCOCF<sub>2</sub>COF (I) reacted with tetrafluoroethylene oxide (II) in diglyme containing CsF at -25° to give mixts. containing MeO<sub>2</sub>CCF<sub>2</sub>(CF<sub>2</sub>OCF<sub>2</sub>)<sub>n</sub>CO<sub>2</sub>Me (n = 1-6) after quenching with MeOH; FCO(CF<sub>2</sub>)<sub>4</sub>COF reacted analogously with II to give MeO<sub>2</sub>C(CF<sub>2</sub>)<sub>4</sub>(CF<sub>2</sub>OCF<sub>2</sub>)<sub>n</sub>CO<sub>2</sub>Me (n = 1-3). Reaction of I with hexafluoropropylene oxide afforded products of addition at both carbonyl groups of I, i.e., MeO<sub>2</sub>C[CF(CF<sub>3</sub>)OCF<sub>2</sub>]<sub>m</sub> CF<sub>2</sub>[CF<sub>2</sub>OCF(CF<sub>3</sub>)]<sub>n</sub>CO<sub>2</sub>Me (m = n = 1,2; m = 1-3, n = m-1).  
 IT 50733-65-8P  
 RL: SPN (Synthetic preparation); PREP (Preparation)  
 (preparation of)  
 RN 50733-65-8 CAPLUS

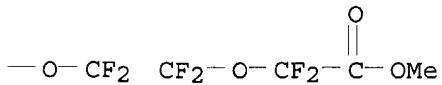
10/630,698

CN 3,6,9,12,15,18-Hexaoxaheneicosanedioic acid, 2,2,4,4,5,5,7,7,8,8,10,10,11,  
11,13,13,14,14,16,16,17,17,19,19,20,20-hexacosafuoro-, dimethyl ester  
(9CI) (CA INDEX NAME)

PAGE 1-A



PAGE 1-B



=> log hold

COST IN U.S. DOLLARS

SINCE FILE ENTRY	TOTAL SESSION
139.32	765.69

FULL ESTIMATED COST

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

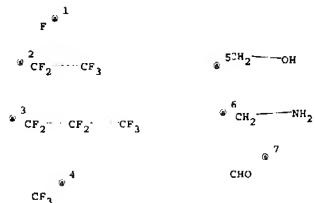
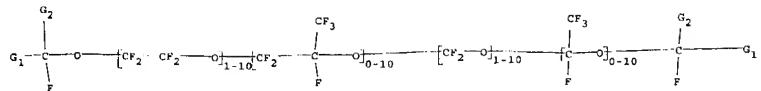
SINCE FILE ENTRY	TOTAL SESSION
-5.60	-14.70

CA SUBSCRIBER PRICE

SESSION WILL BE HELD FOR 60 MINUTES

STN INTERNATIONAL SESSION SUSPENDED AT 07:38:59 ON 10 NOV 2004

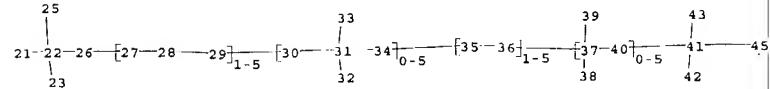
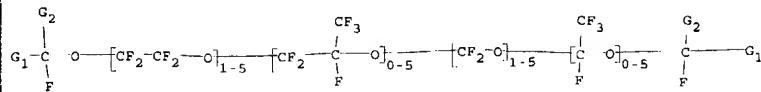
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to search

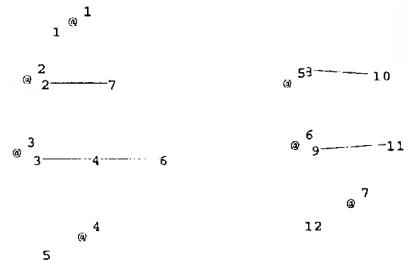
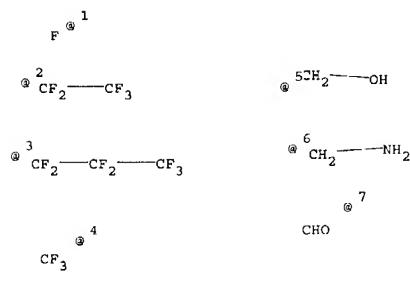
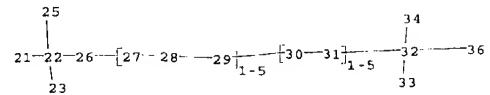
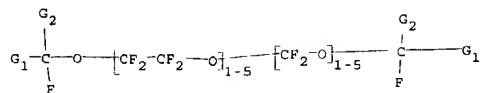


chain nodes :  
 1 2 3 4 5 6 7 8 9 10 11 12 21 22 23 25 26 27 28 29 30 31 32 33  
 34 35 36 37 38 39 40 41 42 43 45  
 chain bonds :  
 2-7 3-4 4-6 8-10 9-11 21-22 22-23 22-25 22-26 26-27 27-28 28-29 29-30 30-31  
 31-32 31-33 31-34 34-35 35-36 36-37 37-40 37-38 37-39 40-41 41-42 41-43  
 41-45  
 exact/norm bonds :  
 21-22 22-25 22-26 31-34 36-37 37-40 40-41 41-43 41-45  
 exact bonds :  
 2-7 3-4 4-6 8-10 9-11 22-23 26-27 27-28 28-29 29-30 30-31 31-32 31-33 34-35  
 35-36 37-38 37-39 41-42  
  
 G1:[\*1],[\*2],[\*3],[\*4],[\*5],[\*6],[\*7]  
 G2:[\*1],[\*2]  
 Match level :  
 1:CLASS 2:CLASS 3:CLASS 4:CLASS 5:CLASS 6:CLASS 7:CLASS 8:CLASS 9:CLASS  
 10:CLASS 11:CLASS 12:CLASS 21:CLASS 22:CLASS 23:CLASS 25:CLASS 26:CLASS 27:CLASS  
 28:CLASS 29:CLASS 30:CLASS 31:CLASS 32:CLASS 33:CLASS 34:CLASS 35:CLASS 36:CLASS  
 37:CLASS 38:CLASS 39:CLASS 40:CLASS 41:CLASS 42:CLASS 43:CLASS 45:CLASS

C:\Program Files\Stnexp\Queries\10630698a.str

Structure too large to search





chain nodes :

1 2 3 4 5 6 7 8 9 10 11 12 21 22 23 25 26 27 28 29 30 31 32 33  
34 36

chain bonds :

2-7 3-4 4-6 8-10 9-11 21-22 22-23 22-25 22-26 26-27 27-28 28-29 29-30 30-31  
31-32 32-36 32-33 32-34

exact/norm bonds :

21-22 22-25 22-26 31-32 32-36 32-34

exact bonds :

2-7 3-4 4-6 8-10 9-11 22-23 26-27 27-28 28-29 29-30 30-31 32-33

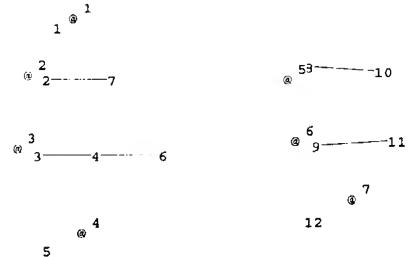
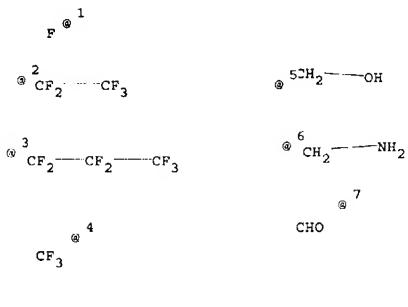
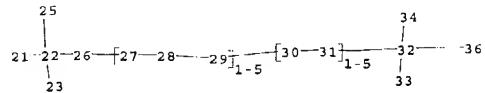
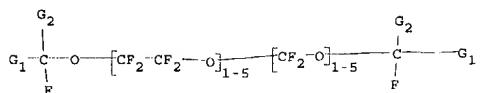
G1:[\*1],[\*2],[\*3],[\*4],[\*5],[\*6],[\*7]

G2:[\*1],[\*2]

Match level :

1:CLASS 2:CLASS 3:CLASS 4:CLASS 5:CLASS 6:CLASS 7:CLASS 8:CLASS 9:CLASS  
10:CLASS 11:CLASS 12:CLASS 21:CLASS 22:CLASS 23:CLASS 25:CLASS 26:CLASS 27:CLASS  
28:CLASS 29:CLASS 30:CLASS 31:CLASS 32:CLASS 33:CLASS 34:CLASS 36:CLASS

C:\Program Files\Stnexp\Queries\10630698b-1.str



chain nodes :

1 2 3 4 5 6 7 8 9 10 11 12 21 22 23 25 26 27 28 29 30 31 32 33  
34 36

chain bonds :

2-7 3-4 4-6 8-10 9-11 21-22 22-23 22-25 22-26 26-27 27-28 28-29 29-30 30-31  
31-32 32-36 32-33 32-34

exact/norm bonds :

21-22 22-25 22-26 31-32 32-36 32-34

exact bonds

**2-7    3-4    4-6    8-10    9-11    22-23    26-27    27-28    28-29    29-30    30-31    32-33**

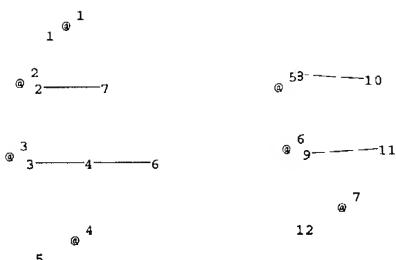
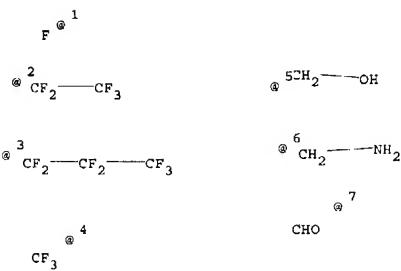
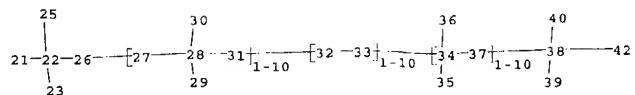
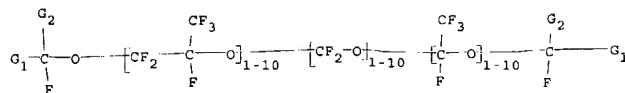
G1:[\*1],[\*2],[\*3],[\*4],[\*5],[\*6],[\*7]

G2: [\*1], [\*2]

Match level :

1:CLASS 2:CLASS 3:CLASS 4:CLASS 5:CLASS 6:CLASS 7:CLASS 8:CLASS 9:CLASS  
10:CLASS 11:CLASS 12:CLASS 21:CLASS 22:CLASS 23:CLASS 25:CLASS 26:CLASS 27:CLASS  
28:CLASS 29:CLASS 30:CLASS 31:CLASS 32:CLASS 33:CLASS 34:CLASS 36:CLASS

Structure too large to  
Search



chain nodes :

1 2 3 4 5 6 7 8 9 10 11 12 21 22 23 25 26 27 28 29 30 31 32 33  
34 35 36 37 38 39 40 42

chain bonds :

2-7 3-4 4-6 8-10 9-11 21-22 22-23 22-25 22-26 26-27 27-28 28-29 28-30 28-31  
31-32 32-33 33-34 34-37 34-35 34-36 37-38 38-39 38-40 38-42

exact/norm bonds :

21-22 22-25 22-26 28-31 33-34 34-37 37-38 38-40 38-42

exact bonds :

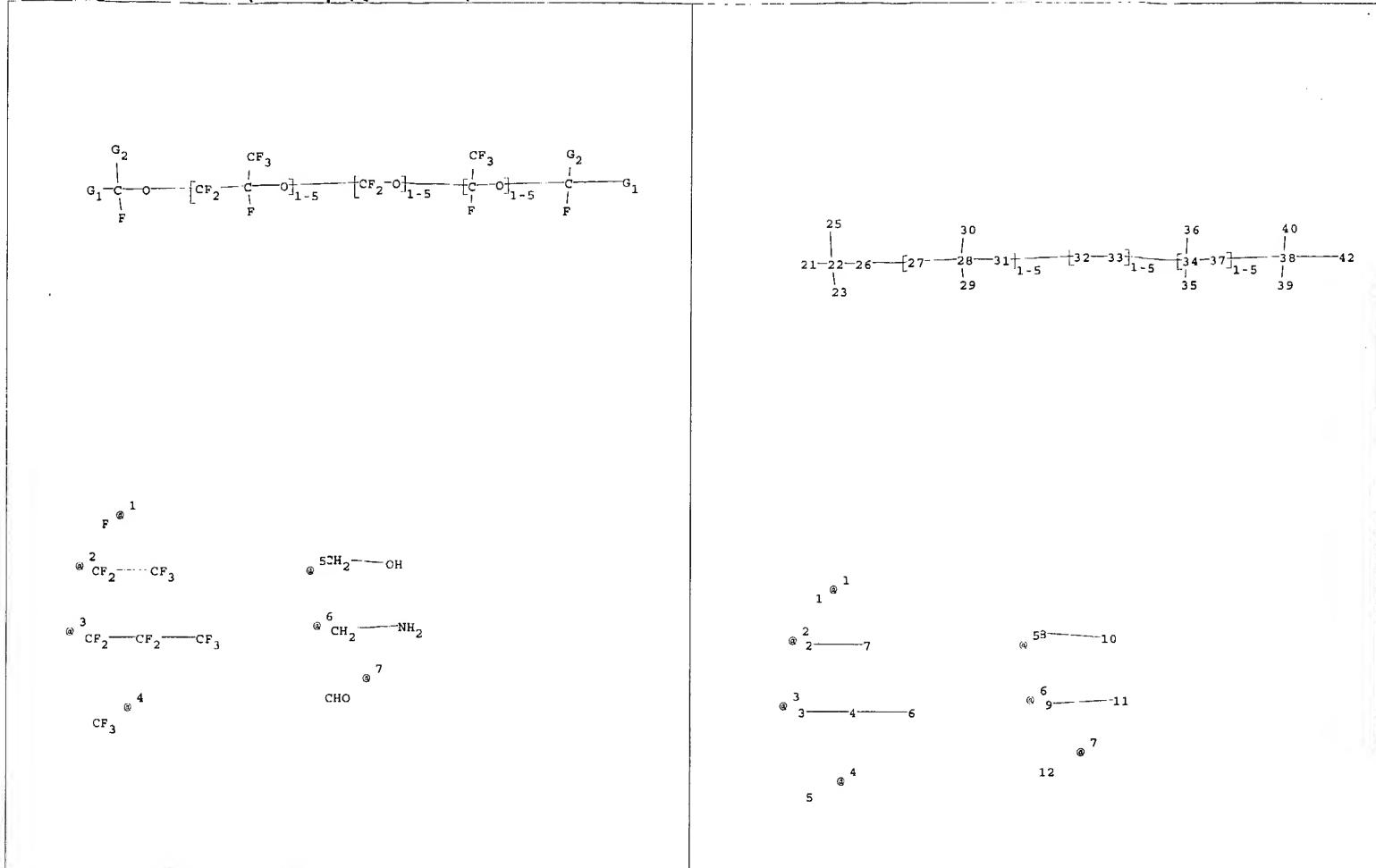
2-7 3-4 4-6 8-10 9-11 22-23 26-27 27-28 28-29 28-30 31-32 32-33 34-35 34-36  
38-39

G1:[\*1],[\*2],[\*3],[\*4],[\*5],[\*6],[\*7]

G2:[\*1],[\*2]

Match level :

1:CLASS 2:CLASS 3:CLASS 4:CLASS 5:CLASS 6:CLASS 7:CLASS 8:CLASS 9:CLASS  
10:CLASS 11:CLASS 12:CLASS 21:CLASS 22:CLASS 23:CLASS 25:CLASS 26:CLASS 27:CLASS  
28:CLASS 29:CLASS 30:CLASS 31:CLASS 32:CLASS 33:CLASS 34:CLASS 35:CLASS 36:CLASS  
37:CLASS 38:CLASS 39:CLASS 40:CLASS 42:CLASS



chain nodes :

1	2	3	4	5	6	7	8	9	10	11	12	21	22	23	25	26	27	28	29	30	31	32	33
34	35	36	37	38	39	40	42																

chain bonds :

2-7	3-4	4-6	8-10	9-11	21-22	22-23	22-25	22-26	26-27	27-28	28-29	28-30	28-31	
31-32	32-33	33-34	34-37	34-35	34-36	37-38	38-39	38-40	38-42					

exact/norm bonds :

21-22	22-25	22-26	28-31	33-34	34-37	37-38	38-40	38-42
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exact bonds :

2-7	3-4	4-6	8-10	9-11	22-23	26-27	27-28	28-29	28-30	31-32	32-33	34-35	34-36
38-39													

G1:[\*1],[\*2],[\*3],[\*4],[\*5],[\*6],[\*7]

G2:[\*1],[\*2]

Match level :

1:CLASS	2:CLASS	3:CLASS	4:CLASS	5:CLASS	6:CLASS	7:CLASS	8:CLASS	9:CLASS
10:CLASS	11:CLASS	12:CLASS	21:CLASS	22:CLASS	23:CLASS	25:CLASS	26:CLASS	27:CLASS
28:CLASS	29:CLASS	30:CLASS	31:CLASS	32:CLASS	33:CLASS	34:CLASS	35:CLASS	36:CLASS
37:CLASS	38:CLASS	39:CLASS	40:CLASS	42:CLASS				